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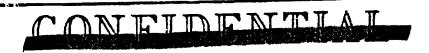
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EIGHTH QUARTERLY PROGRESS REPORT

ON

DISSEMINATION OF SOLID AND LIQUID BW AGENTS

(Unclassified Title)



For Period March 4, 1962 - June 4, 1962 Contract No. DA-18-064-CML-2745

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Project No: 82408
Date: August 22, 1962

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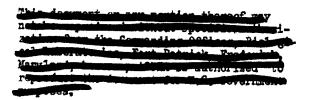
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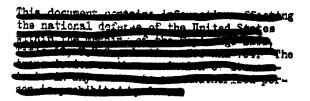
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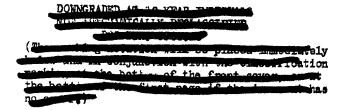
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#### **FOREWORD**

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#### ABSTRACT

This Eighth Quarterly Progress Report covers the work accomplished during the last reporting period on research and development related to the dissemination of BW agents.

The progress in theoretical and experimental studies of the mechanics of dry powders is reported. This work continues to produce basic information which is being applied in the design and development of an airborne disseminator.

Operation of the aerophilometer, with aerosols of dry powders, for the purpose of studying experimental techniques and determining operating parameters is discussed. This preliminary work is essential to carrying out the program on stability properties of aerosols.

Data on the effects of compaction and additives on the viability of <u>Sm</u> powder are presented.

The tests at Fort Detrick using the GMI-3 fixture and wind tunnel to generate aerosols in the 40-foot test sphere are described.

Results with the full-scale experimental equipment for feeding and metering compacted dry powders are reported.

The preliminary design of an airborne dry BW agent disseminating store is presented and described.

Successful flight tests of the liquid BW agent disseminating store on an A4D-1 airplane are discussed.

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EIGHTH QUARTERLY PROGRESS REPORT 2019

### ON DISSEMINATION OF SOLID AND LIQUID BW AGENTS

#### 1. INTRODUCTION

This is the Eighth Quarterly Progress Report on work accomplished on Contract Number DA-18-064-CML-2745. Under this contract, General Mills, Inc. is conducting a comprehensive research and development program on the dissemination of solid and liquid BW agents.

With the completion of the developmental model of the liquid agent disseminating store in April of this reporting period, the emphasis of work shifted almost entirely to areas relating to finely-divided solids. Consequently most of this report deals with progress in these areas. However, Section 8 does present some of the results of laboratory and field testing of the liquid agent disseminator.

On 22, 23 May, General Mills, Inc. was host to the Third Dissemination Coordination Meeting which was attended by representatives from Fort Detrick, Army Chemical Center, Chemical Corps Research and Development Command, Aerojet-General Corporation, Cornell Aeronautical Laboratories and General Mills, Inc. Some of the material covered in this progress report was also presented at this meeting.

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## 2. THEORETICAL AND EXPERIMENTAL STUDIES OF THE MECHANICS OF DRY POWDERS

Studies of the mechanical properties and behavior of dry powders were continued during the period covered by this report. Experimental results on the compaction characteristics, shear strength, and tensile strength of various powders are presented.

#### 2.1 Compaction Experiments with Particulate Media

Results obtained with an improved version of the piston-cylinder compaction apparatus and a hydrostatic compaction device are discussed and compared in the following sections.

#### 2.1.1 Piston-Cylinder Compaction Experiments

As pointed out in the last quarterly report (seventh), the initial piston-cylinder apparatus used in the compaction experiments was subject to several limitations and, consequently, an improved piece of apparatus (Figure 2.1\*) was fabricated. Powder is placed in the cylinder at the base of the apparatus. A surface stress is applied to the powder by a piston which is lowered by turning the crank at the top of the apparatus. An aluminum ring equipped with strain gages is used to measure the stress. Large-scale movement of the piston is measured by means of a Starrett dial indicator, and small-scale movement is measured by means of a sensitive differential transformer. The output of the strain gages and the differential transformer is recorded on a Sanborn recorder.

Using the experimental results obtained from the tests employing this piece of apparatus, a relationship between the bulk density of the powder and the stress necessary to obtain this density was obtained. These results

<sup>\*</sup> This is the same apparatus shown in the referenced report.

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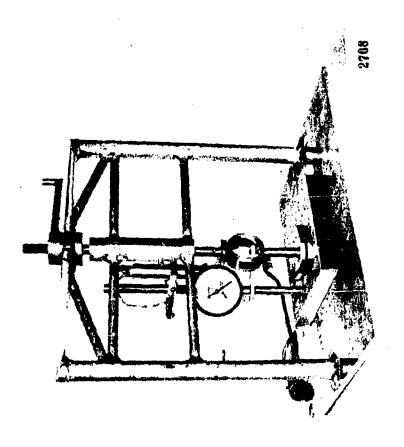


Figure 2. 1 Improved Piston-Cylinder Compaction Apparatus

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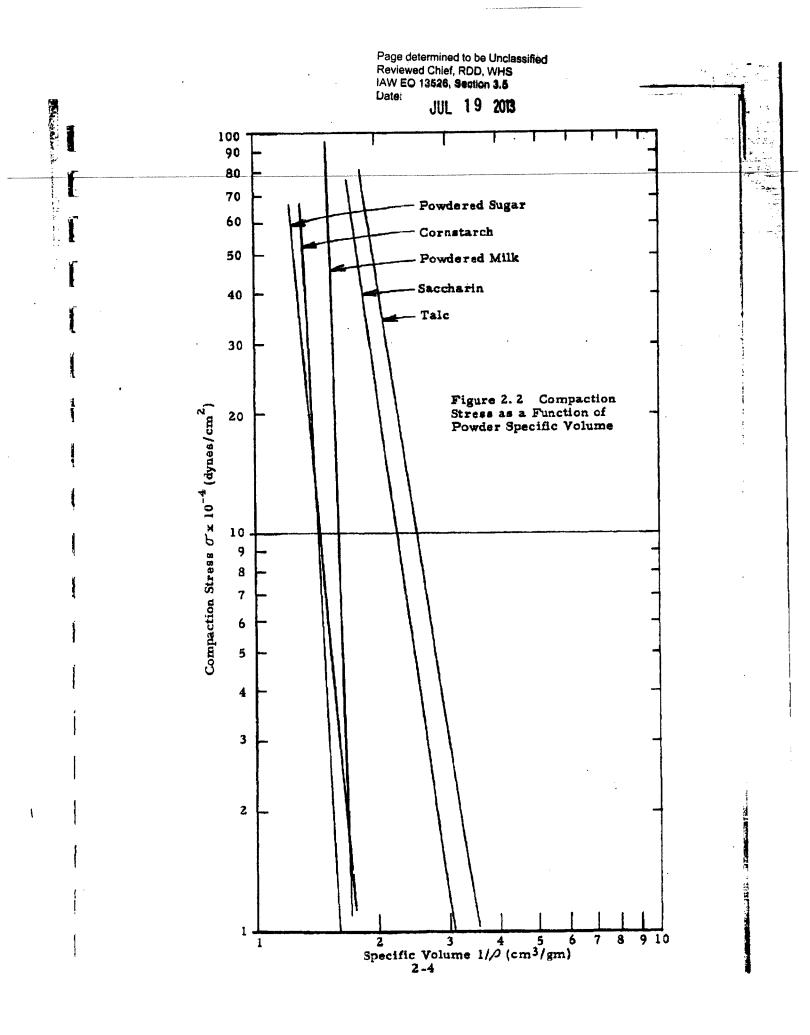
are plotted in Figure 2.2 for several different powders. The stress levels for these tests were considerably higher than those of earlier tests. The stress was increased monotonically during the tests.

A plot of the stress versus reciprocal density is a straight line on log-log paper which is in agreement with the previous results. An empirical formula of the form  $\mathcal{O} = k(1/\mathcal{O})^T$  can be used as a good fit to the data. The constants of this equation are given below for the powders tested:

Powder	r	k	
Talc	- 6.50	4.42 × 10 <sup>7</sup>	
Saccharin	- 7.70	2. 21 × 10 <sup>7</sup>	
Powdered Sugar	-11.5	6.8 × 10 <sup>6</sup>	
Powdered Milk	-35. 9	$1.6 \times 10^9$	
Cornstarch	-20.8	1.67 × 10 <sup>8</sup>	

At each compaction state a certain amount of energy is stored in the powder bed in the form of elastic energy as pointed out in an earlier report. In the initial tests it was assumed that the material had linear elastic characteristics. It is now necessary to revise this assumption on the basis of more accurate measurements carried out with the improved apparatus. The general shape of the elastic recovery versus stress curve for powders such as tale, saccharin, and powdered sugar is shown in Figure 2.3.

It is interesting to note that very little energy is recovered as the stress is reduced at high stress levels, but that most of the elastic recovery takes place at low stress levels. This implies that considerably less elastic energy is stored in the powder bed for a particular compaction state than was originally assumed.



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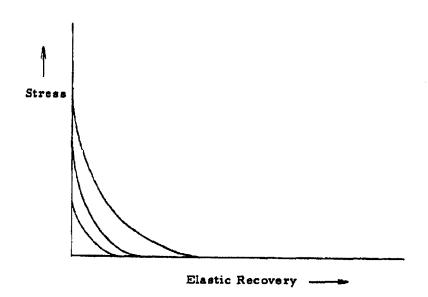


Figure 2.3 Shape of Elastic Recovery versus Stress Curve

Some difficulty was experienced in obtaining precise measurements of the elastic behavior of powder due to insufficient rigidity of the new compaction energy apparatus. At high stress levels the apparatus had a tendency to flex, making precise measurements of the elastic recovery of a powder difficult to obtain. However, the basic nature of the elastic recovery, illustrated in Figure 2.3, seems to be definitely established. In order to obtain more precise data, it will be necessary to modify the apparatus to make it more rigid.

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In the process of testing the various powders it was noticed that powdered milk and cornstarch behaved in a unique manner during the compaction process. As the loading piston was gradually advanced, the stress was found to increase rapidly and then fall sharply to a lower level. In other words the powder seemed to compact in a stick-slip fashion. Figure 2.4 is a comparison of the recording traces showing the variation of stress as a function of time for tale and powdered milk as the powders were gradually compacted.

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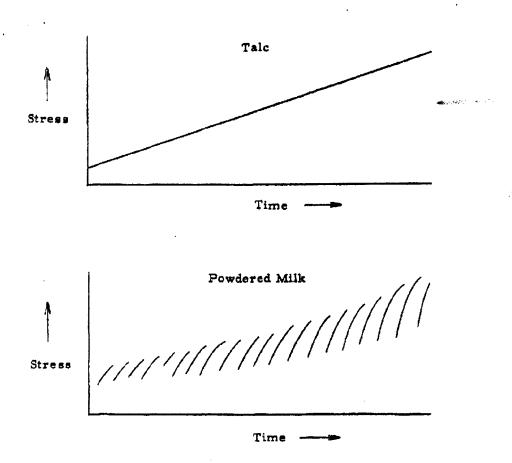


Figure 2.4 Compaction Behavior of Talc and Powdered Milk

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The stick-slip phenomena observed with powdered milk and cornstarch diminishes at high stress levels. The stresses plotted in Figure 2.2 for powdered milk and cornstarch were the values noted after the stress had fallen from the peak value. The magnitude of the stress peaks was influenced by the rate at which a force was applied. If the force was applied rapidly the peaks were quite small. This phenomenon, indicating that the rate at which surface stresses are applied may be an important factor in the compaction process, will be investigated in future tests.

An expression for the compaction energy as a function of density can be obtained from the stress-density curve (Figure 2.2) and is of the form  $W = C \rho^n$ . However, from the previous discussion it is apparent that the stress necessary to reach a given compaction state, and hence the energy involved in the process, may depend on the path taken to reach that compaction state. For example, if a certain stress is applied and released several times in compacting a bed of powder, the energy of compaction might be different than if the stress were applied monotonically to reach the same compaction state. In those tests made to date in which the test procedure was varied, the results have not been significantly different. However, an investigation to precisely determine the effect of path on the compaction energy is planned for future work.

#### 2, 1, 2 Hydrostatic Compaction Experiments

In the process of performing the piston-cylinder compaction experiments, the question arose as to how much the results depended upon the constraints and loading conditions imposed by the apparatus. In order to answer this question a hydrostatic compaction experiment was devised in which the constraints imposed by the apparatus were minimized. The apparatus is shown in Figure 2.5.

A sample of powder was put into a polyethylene bag which was then hermetically sealed. The bag was inserted in the pressure chamber with the powder sample vented to the outside. The chamber was then filled with

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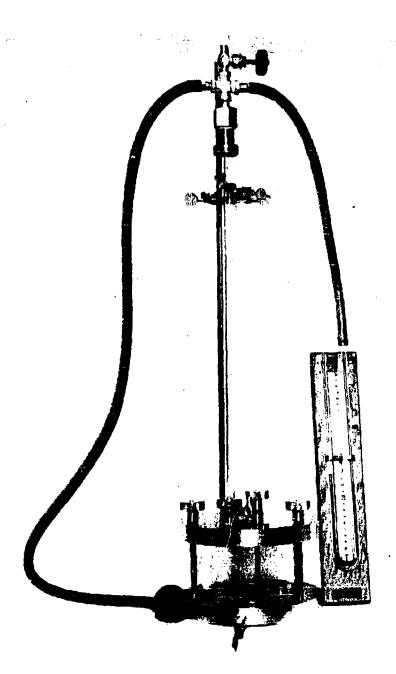


Figure 2.5 Hydrostatic Compaction Apparatus

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water to a certain level in the calibrated tube used for measuring volume change. Then the pressure in the chamber was increased in small increments and the volume change was noted. It was interesting to note that considerable time was required for the system to come to equilibrium after the pressure was increased. Evidently the stress distribution in a powder bed undergoes a change during the compaction process. After the system reached a state of equilibrium, the hydrostatic pressure was released and the volume change caused by elastic recovery was noted. The shape of the elastic recovery versus stress curves was essentially the same as that shown in Figure 2. 3.

From the data, a curve of stress versus the reciprocal density was plotted for the two powders tested (tale and saccharin) and a straight line on log-log paper was obtained. Thus, both the hydrostatic and piston-cylinder compaction tests yield a stress-density relationship which is described by the empirical formula:

$$\sigma = k \left(\frac{1}{\rho}\right)^{x}.$$

The constants for this equation for the two powders tested in the hydrostatic compaction apparatus are:

talc: 
$$k = 1.39 \times 10^6$$
,  $r = -7.6$ ;

and

saccharin: 
$$k = 1.95 \times 10^7$$
,  $r = -6.24$ .

Values of the exponent r are in good agreement with those determined by the piston-cylinder method. The coefficient k for saccharin is also in good agreement with the results of earlier tests. However, the value of k for talc differed appreciably from that determined by the piston-cylinder method. This apparent discrepancy might be attributable to differences in

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the compaction process. At the conclusion of one of the hydrostatic experiments with talc, the stress was cycled several times. The result of this cycling caused the bulk density to increase with each succeeding cycle until a point was reached where the density did not seem to increase further. From this one can see that it is somewhat ambiguous to speak of the stress necessary to reach a certain compaction state without specifying the compaction process, at least for a powder like talc. Other powders may respond differently to the cycling process.

The hydrostatic compaction apparatus has several drawbacks, one being expansion of the chamber under pressure. This expansion necessitated calibration tests, resulting in a tare which has to be taken into account to determine the true volume change of the powder sample for various pressures. Some difficulty was also experienced in determining the exact volume of the powder sample, a value needed in order to calculate the bulk density. Nevertheless, it is believed that the results are meaningful and are in accord with our previous results.

In view of the fact that results from hydrostatic compaction tests seem to be in close agreement with the results obtained from the piston-cylinder tests, future compaction tests will be conducted utilizing the piston-cylinder arrangement because of the relative ease of conducting these tests.

The powders that have been investigated thus far seem to fall into two major groups. In the first group are powders such as powdered milk and cornstarch which display the above-mentioned stick-slip phenomena and compact very little even when subjected to high compressive stresses. In the second group are powders such as talc and saccharin which compact much more smoothly.

Future tests will be oriented to determine what factors are affecting the compacting characteristics of a powder. Factors such as the history of the powder, particle shape and size, electrostatic charge, and moisture content would seem to be among the most important factors to consider.

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#### 2. 2 Powder Shear Strength Studies by the Triaxial Test Technique

Preliminary tests with the triaxial test apparatus described in a previous report revealed deficiencies in the originally proposed test technique, which produced excessive scatter of the experimental data. To improve the reliability and reproducibility of the triaxial shear test, modifications were made in the method of sample preparation and in the means used to apply stresses to the test sample.

#### 2.2.1 Sample Preparation

For incompactable materials, which exhibit dilatancy in shear, the preparation of homogeneous test samples is a straightforward matter. However, in the case of compactable materials, it does not appear to be possible to produce truly homogeneous, isotropic samples. Since the test sample has the form of a right circular cylinder, it is natural to compact the sample to the degree necessary for handling by employing a pistoncylinder apparatus. However, from prior study of the compaction process in a piston-cylinder device<sup>3</sup>, it is known that the compressive stresses applied to the powder decay exponentially with distance from the loading piston. Furthermore, the ratio of radial to axial stress appears to lie in the range 0, 3 to 0.5 for most materials. From this it may be inferred that the density of the sample will decrease with distance from the loading piston and that the powder will compact non-isotropically. It is clear that the latter effect is not objectionable since it is typical of the behavior of a compactable powder. Non-uniform sample density is undesirable, however, since the shear strength of a powder will, in general, depend on the density.

Several techniques have been used to prepare samples for the triaxial shear test. The technique originally used was based upon the observation that a region of nearly constant density exists in a column of powder which

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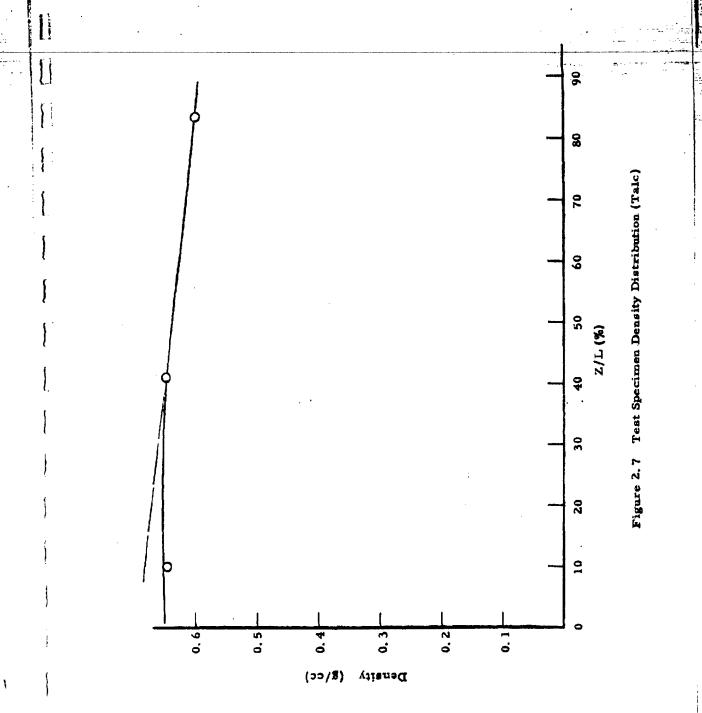
has been compacted in a vertically disposed piston-cylinder device. It was thought that a test sample obtained from this region would be ideal for tri-axial testing. However, it was found to be impossible to obtain a sufficient degree of compaction in this region to permit the minimal handling required for preparation of the test specimen. Thus, it became evident that a density gradient in the sample would have to be accepted for samples compacted in the piston-cylinder device. A series of experiments were carried out to determine the density variation along a powder column for loads sufficiently high to cause compaction to occur along the entire column. Typical results for talc are shown in Figure 2.6 for two applied loads. It is apparent that the density gradient is essentially constant over the length of the compacted powder column.

This result led to a modification of our sample preparation technique whereby a known mass of powder is compacted to a preselected final length, thus precisely defining the mean sample density. This technique has been found to yield improved control over sample uniformity, as judged by the reproducibility of shear test results obtained by using samples prepared in this way.

Actually, this method of preparing test specimens is very similar to the proposed technique described in the last quarterly report. At high compaction stresses, however, it was found that external support was required for the split tube within which the powder sample is confined during compaction. Another essential change was to roughen the inner surfaces of the split tube to prevent stretching of the rubber membrane during compaction.

The variation of density with axial distance from the loading piston for the modified sample preparation technique is shown in Figure 2.7. The drop in density near the high-density end of the sample appears to be caused by elastic relaxation effects. (See Figure 2.6 also.)

Figure 2. 6 Density Variation in a Compacted Powder Column (Talc)



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#### 2. 2. 2 Theoretical Density Variation

The density variation in a powder sample which has been compacted in a piston-cylinder device may be determined analytically by combining results from the piston-cylinder theory with the empirical relationship between density and applied stress 2, 3:

$$\sigma = C\rho^{m} \tag{2.1}$$

From the piston-cylinder theory, we have:

$$C'(Z) = \sigma(0) e^{-\frac{4KL}{D}} (\frac{Z}{L})$$
 (2.2)

Eliminating the axial stress  $\sigma(Z)$ , Equations (2.1) and (2.2) lead to the relationship:

$$\rho(Z) = \rho(0) e^{-\frac{4KL}{mD}} (\frac{Z}{L})$$
 (2.3)

In this equation the wall resistance parameter K depends on the shear angle \$\textit{\theta}\$ of the powder and the wall friction angle \$\theta\$, while m is an empirically determined powder parameter.

Computed values of  $\rho(Z)$  from Equation (2.3) are shown in Figure 2.6 for  $\sigma(0) = 14.6 \times 10^6$  dynes/cm<sup>2</sup>, m = 5.46 (Reference 3) and for two values of the wall resistance parameter: K = 0.358 and K = 0.200. The value K = 0.358 (corresponding to the maximum theoretical value of the wall resistance parameter<sup>2</sup>) yields the maximum theoretical rate of decay of density. The smaller value of K = 0.200 was obtained from Figure 2.1.3, Reference 2, for the experimental values:  $\beta = 40^\circ$  (talc),  $0 = 35^\circ$  (talc on aluminum). The theoretical curve for K = 0.200 agrees rather well with the experimental curve for  $\sigma(0) = 14.6 \times 10^6$  dynes/cm<sup>2</sup>.

#### 2. 2. 3 Test Procedure and Typical Test Results

During initial work with the triaxial test technique, it was found that the stress at which yielding of the sample occurred was difficult to define. The principal reason for this appears to be the constraining effect of the rubber membrane, which prevents a sudden collapse of the powder when shear failure occurs. In an effort to increase the precision of the test and to secure a better indication of sample failure, the triaxial test apparatus was adapted for use with an available Instron test machine. Through use of the Instron, the following desirable features were achieved: 1) accurate force measurement through use of a sensitive load cell, 2) precise and adjustable rate of strain, and 3) accurate recording of applied axial load as a function of sample strain.

Figure 2.8 shows the triaxial test apparatus mounted in the Instron test machine. The test procedure is as follows:

- The test specimen, prepared in the manner described above, is installed in the pressure chamber of the test apparatus and properly centered (locating pins are provided for this purpose in the sample end fittings).
- The test chamber is sealed and pressured to the desired chamber pressure.
- After checking to make sure there is no leakage to the sample interior, the machine is started and run until shear failure occurs.

In these tests, shear failure is evidenced by a sudden change in the slope of the load-strain curve. This is illustrated by Figure 2.9 showing a series of load-strain curves for a mean sample density of 0.650 g/cc and a chamber pressure of 1 psi. The appearance of the test specimens after failure is shown in Figure 2.10. It is apparent from the photograph that failure occurs by the formation of cracks in the compacted powder, although the rubber membrane prevents collapse of the sample.

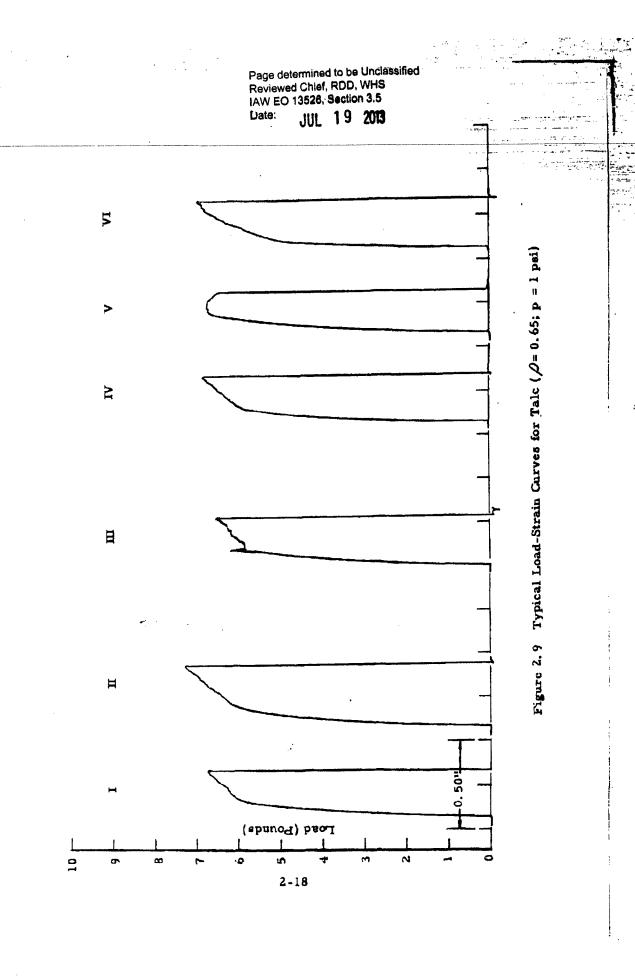
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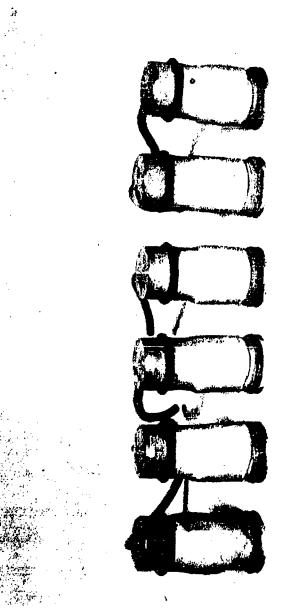
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Figure 2.8 Triaxial Test Apparatus



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Test Specimens Illustrating Shear Failure (O=0.65; p = 1 psi) Figure 2, 10

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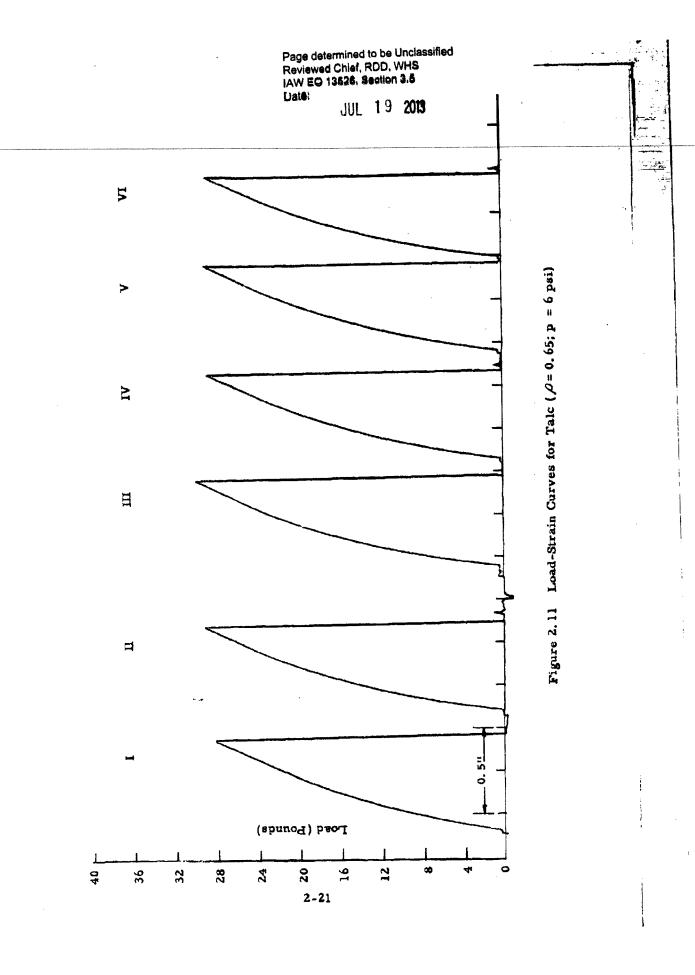
The results shown in Figures 2.9 and 2.10 are typical of runs carried out for pressures in the range 0 to 2 psi. At higher pressures, however, it was impossible to detect sample failure since the sample merely bulged outward without a break in the load-strain curve. This behavior is illustrated by Figures 2.11 and 2.12. It is suspected that this behavior may be caused by additional compaction during the test since the applied loads are comparable to the loads used in preparing the sample. Of course, the effect of the membrane in this case is to stabilize the powder as bulging occurs. There is little reason to doubt that the high loads shown in Figure 2.11 for a chamber pressure of 6 psi are attainable only because of the stabilizing

The yield locus established from tests carried out at pressures of 0, 0.24, 0.50 and 2.0 psi is plotted in Figure 2.13. The slope of the yield locus is about 35 degrees, which agrees quite well with the shear angle determined from sliding disk measurements  $^2$  ( $\theta = 40$  degrees).

effect of the membrane.

The experiments and test results reported herein have been primarily concerned with the development of techniques for application of the triaxial test to compactable powders. Additional tests are planned for future work in which powders other than talc will be tested and the effects of such factors as moisture content will be investigated.

Our studies of the triaxial test have disclosed several limitations of this method of conducting shear tests. The most serious limitation is that the test technique is restricted to samples which have been compacted to high densities; also, no means have been found to produce samples of uniform density. Another serious problem is the constraining effect of the rubber membrane, which prevents a "natural" shear failure. It should be noted, however, that the membrane is not required for tests at zero pressure. A further difficulty with the triaxial technique is the difficulty of defining the shear locus for small compressive stresses. Thus the strength of the powder in pure shear cannot be obtained directly from this type of test.



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Figure 2.12 Test Specimens After High-Pressure Tests (Q = 0.65; p = 6 psi)

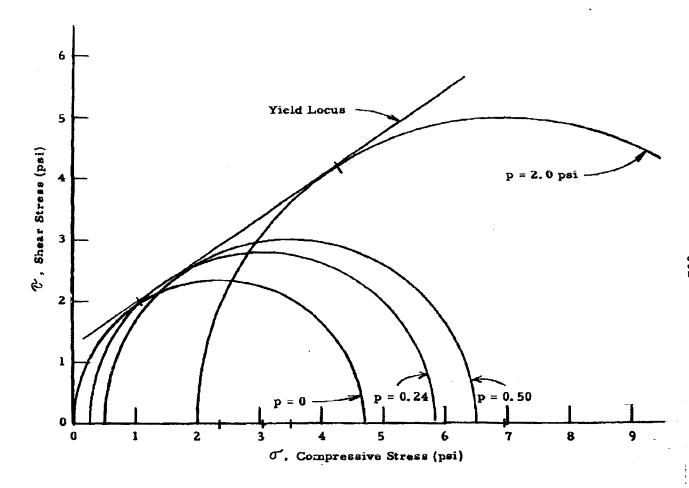


Figure 2.13 Triaxial Test Results for Talc at a Mean Sample Density,  $\rho = 0.65$ 

In spite of these deficiencies, the triaxial test provides much worthwhile information on the behavior of dry powders. Also, it is quite possible that some of the apparent limitations of the triaxial test can be removed through improvements in test techniques.

#### 2.3 Bulk Tensile Strength of Compressed Powders

A new apparatus and technique for the measurement of bulk tensile strength of compressed powders was described in the last quarterly report. Work in recent months has been devoted to an exhaustive study to refine techniques and to establish the most accurate and efficient method for a comparative measure of bulk tensile strength of various bulk powders.

According to theory<sup>2, 4</sup> the bulk tensile strength of a compressed powder is an exponential function of the distance from the compressive piston to the fracture plane:

$$\sigma = \sigma_0 e^{-kL} \qquad (2.4)$$

where:

or = bulk tensile strength of a column of compressed powder at distance L from the piston

= bulk tensile strength of the compressed powder immediately below the piston

k = constant

L = distance from piston to fracture plane.

The method employed involves the compression of bulk powder in a vertical segmented column. The design of the apparatus permits the powder to be fractured by a measured vertical force at several points down the column, permitting the bulk tensile strength to be measured as a function of:

1) bulk density, 2) distance from compressive force application to fracture plane, and 3) total column length.

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Although the bulk tensile strength does vary with length of time of application of compressive load, our recent studies have led to the adoption of a 1-1/2 hour compression time. It is believed that this time period not only is adequate for good reproducibility but also allows the operator to make four efficient runs per working day. In addition to the adoption of 1-1/2 hours as a standard time of compression, three compressive loads (5.55 x  $10^5$ , 7.88 x  $10^5$ , and  $10.59 \times 10^5$  dynes/cm<sup>2</sup>) have been adopted as standard. Although this test as described does not exhaustively cover all possibilities, we believe that it does offer an acceptable method for a comparative study of a physical phenomena which is at best difficult to measure.

Figures 2.14, 2.15 and 2.16 represent typical plots for zinc cadmium sulfide at three compressive loads. The individual plots were made by the method of least squares. Individual points were omitted to afford greater ease of visual comparison of the various plots.

These data show as indicated in the last quarterly report that  $\sigma_0$  is in fact a function of total plug length  $L_t$ . Figure 2.17 shows the relationship between  $\sigma_0$  and  $L_t$  at the three compressive loads. Since the values obtained for  $\sigma_0$  are for a fairly narrow range of values for  $L_t$  we must extend this range before making valid generalizations. The individual plots are however interestingly close to being linear.

In the near future we plan to obtain more data on the variation of  $\sigma_0$  with  $L_t$  for zinc cadmium sulfide and to utilize our standard method to study the bulk tensile strength characteristics of such powders as tale, powdered sugar, powdered milk, saccharin, and cornstarch.

#### 2.4 Frictional Forces Between Powders and Plated Metal Surfaces

A modification of a test method previously described<sup>5</sup> was used to study the frictional force between talc and samples of cold rolled steel that had been electroplated with nickel, cadmium or zinc. In principle the method

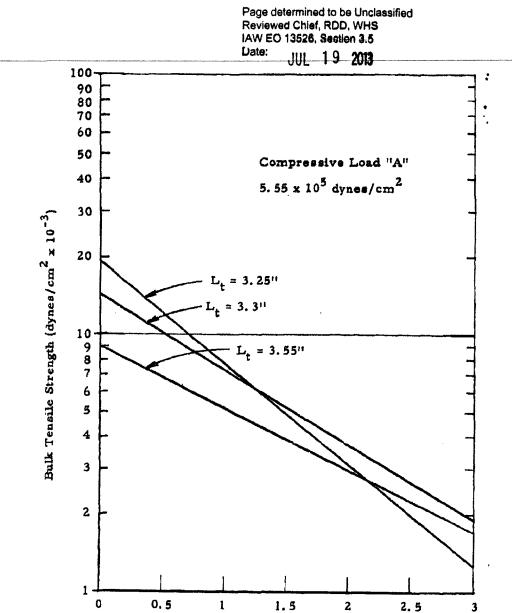


Figure 2.14 Bulk Tensile Strength for Zinc Cadmium Sulfide as a Function of Distance "L" from Compressive Force at Various Total Plug Lengths, L<sub>t</sub>

L (inches)

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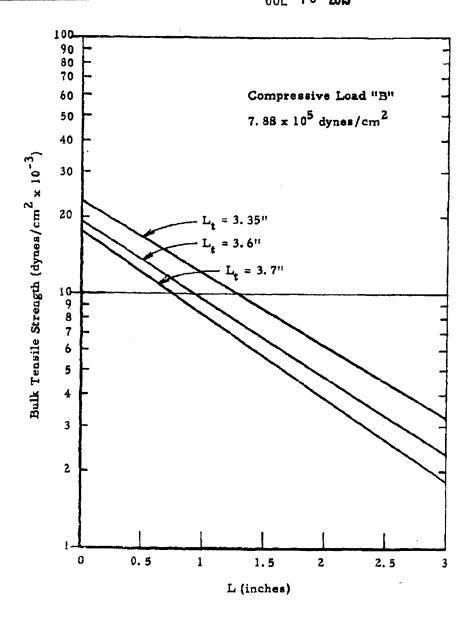


Figure 2.15 Bulk Tensile Strength for Zinc Cadmium Sulfide as a Function of Distance "L" from Compressive Force at Various Total Plug Lengths, L<sub>t</sub>

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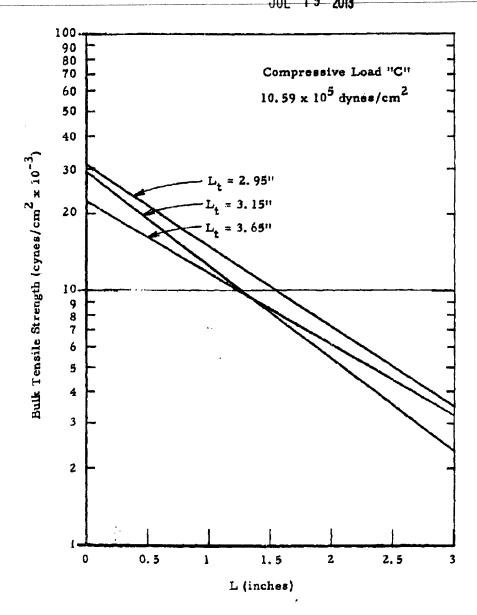


Figure 2.16 Bulk Tensile Strength for Zinc Cadmium Sulfide as a Function of Distance "L" from Compressive Force at Various Total Plug Lengths, L<sub>t</sub>

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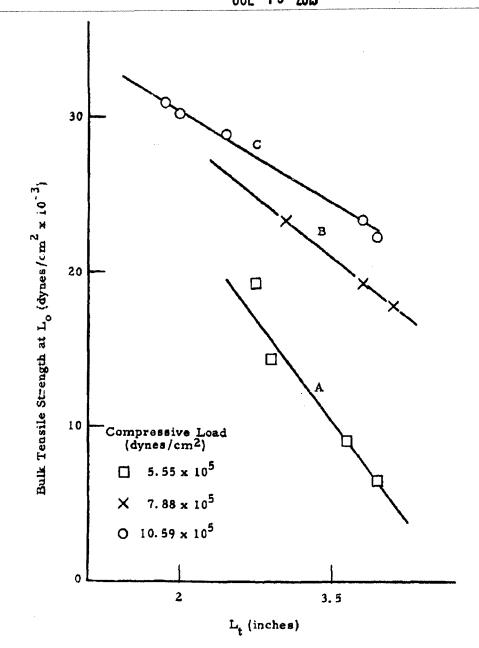


Figure 2.17 Bulk Tensile Strength at L for Zinc Cadmium Solfide as a Function of Total Plug Length L, at Various Compressive Loads

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involves placing a powder on the metal surface and tilting the plate of metal slowly until the powder begins to slide. The slide angle is related to the coefficient of friction as follows:

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where:

# = coefficient of friction

9 = angle of slide

To make a comprehensive study of this problem, the following three methods were used to place the talc upon the metal surface:

- 1) An aluminum ring of 3.5 cm ID was placed on a mechanical vibrator and filled with 1.5 g talc. After a standard 3-minute vibration to assure uniformity of packing, the sample was transferred to the plated metal surface on the tilting table.
- 2) The talc was processed identically to Part (1) except that prior to transfer to the plated surface, the talc was compacted into a plug by the use of a piston assembly with a 9-pound compressive load.
- 3) The tale was processed identically to Part (1) except that after transfer to the plated surface, the tale was compacted directly onto the plated surface by the 9-pound piston assembly.

The tests were performed in a controlled humidity environment (15 to 18 percent relative humidity). Upon analysis of these data, Method (3) was chosen for further study at a higher (>60% RH) humidity. The following is a tabulation of values of angle of slide; each value reported is an average of at least five independent runs:

		Plating			
	Method	Nickel	Cadmium	Zinc	Unplated
Low Humidity (15-18% RH)	1 2 3	40.7 36.0 41.9	42. 6 37. 6 43. 4	42.5 37.8 43.7	43.0 39.7 44.2
High Humidity	3	39. 8	40.0	42.9	42.0

Although Method (2) tends to yield a lower slide angle than Methods (1) or (3), the data indicate only very minor differences in the frictional forces between the talc powder and the various electroplated surfaces. It is interesting to note, however, that regardless of the method utilized, the nickel plate gave consistently the lowest coefficient of friction and the unplated material consistently the highest. It should also be mentioned that in every case when the metal surface had been exposed for some time prior to test, the first test performed on each surface gave values 50 to 75 percent higher than the average. These values were not included in the tabulated data presented. Apparently either surface moisture and/or surface oxidation changes the frictional characteristics of the surface in a way that is removed by making a single test run.

Since there seems to be no obvious advantage in electroplating the metal surfaces of a disseminator in order to reduce frictional forces, no future work in this area is currently planned.

## 2.5 Bulk Density of Compressed Powders

To expand our knowledge of the variation of bulk density in a column of compressed powder, a new apparatus (as shown in Figures 2.18, 2.19 and 2.20) has been designed and perfected. Saccharin has been used as a test powder in our initial series of experiments with the new apparatus.

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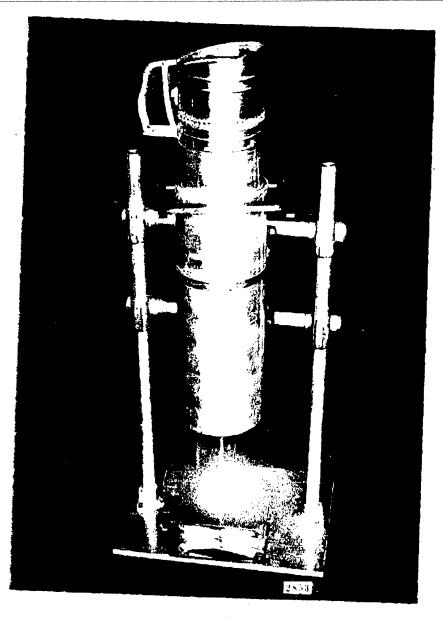


Figure 2, 18 Bulk Density Apparatus Asserobled for Filling

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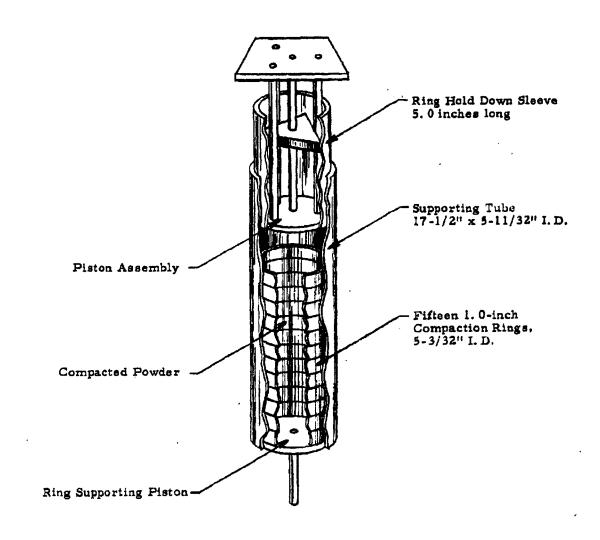


Figure 2.19 Bulk Density Apparatus Assembled for Compaction of Bulk Powders

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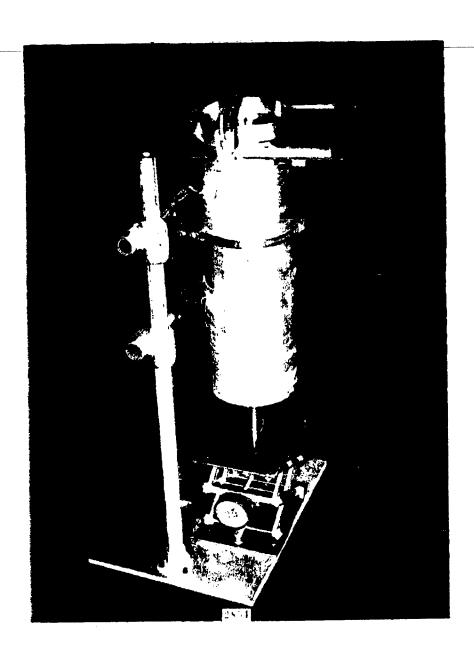


Figure 2, 20 Bulk Density Apparatus Assembled for Cutting of Individual Powder Segments

### 2.5.1 Apparatus and Technique

In an actual experiment the apparatus is assembled as shown in Figure 2.19 but with the sifter replacing the piston assembly (Figure 2.18). The sifter with a mechanical, doorbell-type vibrator attached is intended to aid in uniformity of packing. The powder under study is added slowly through the sifter until the column is filled to the top of the hold-down sleeve. The piston assembly is then put into place and the compressive load applied for a period of one-half hour. The compressive load, piston assembly, and hold-down sleeve are then removed. The entire column of compacted powder plus the compaction rings are moved upward with the aid of the lab jack and ring-supporting piston, exposing the metal rings containing the compacted powder.

The cutting assembly (Figure 2. 20) is then used to divide the plug of compressed powder into one-inch segments of known cross-sectional area. By obtaining the weight of these segments, the bulk density may then be calculated. To obtain further information a "cookie-cutter" assembly is then used to remove a circular portion, representing one-half the cross-sectional area, from the center of each segment. This affords a gross comparison of the variation of bulk density along the radius of the column of compressed powder.

#### 2.5.2 Discussion of Data

The results of an experiment using saccharin as a test powder are shown in Figure 2.21. The compactions were made in order of increasing compressive load with no special attention given to using virgin saccharin each time. To study the possible effects of previous history of compaction of a powder upon bulk density, the 18.9 lb test was repeated using the saccharin that had been used throughout several tests including the 354 lb compression.

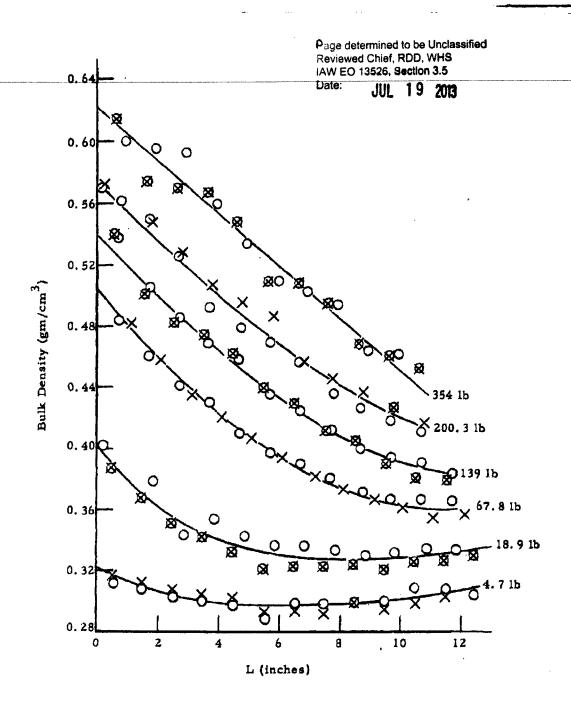


Figure 2.21 Bulk Density of Precompacted Saccharinas a Function of Distance from Piston at Various Compressive Loads

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The two curves are shown for comparative purposes in Figure 2.22. It is interesting to note that the general shapes of the plots are nearly identical. The bulk density of the virgin powder for the first 18.9 lb test was 0.293 gm/cm<sup>3</sup> compared with 0.332 gm/cm<sup>3</sup> for the previously compacted saccharin used in the last 18.9 lb test. These experiments accentuate the dependence of bulk density upon the previous compaction history of the sample and the need for the use of virgin materials for each test in order to obtain reliable comparative results. With this background of experience the experiment was then repeated using virgin saccharin for each run. The results shown in Figure 2.23 indicate a linear relationship between the bulk density and distance from compressive piston. The deviation from linearity at greater distances from the piston will be included in the discussion of Figure 2.25.

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As stated earlier in this report, data were obtained concerning the variation in bulk density along the radius of the plug. Since this variation was so uniform throughout each test, the data shown in Figure 2.24 are presented as typical of the variation found. In order to determine whether or not the radial variation in bulk density is due to the effects of the applied compressive load or to the method of filling the apparatus, an experiment was just completed at the writing of this report in which the bulk density apparatus was filled with saccharin in a manner identical to previous experiments and the bulk density measured without the application of a compressive load. The results are shown in Figure 2.25. Although the apparatus permitted accurate measurement of the average bulk density of the segments, only near the bottom of the column was the powder plug rigid enough to obtain data on the inner and outer segments. It is very interesting to note, however, that radial variation in bulk density of the order of magnitude observed with the compacted powder is also in evidence here. The radial variation is thus affected to a considerable extent by the method of filling the column. It is believed, however, that the vertical variation of bulk density should be relatively unaffected by this phenomena.

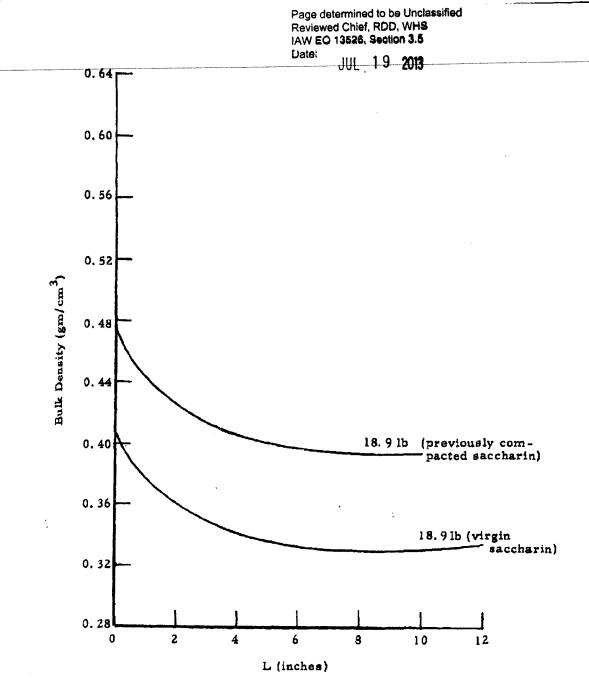


Figure 2. 22 Bulk Density of Saccharin as a Function of Distance from Piston (A comparison of virgin saccharin with previously compacted saccharin.)

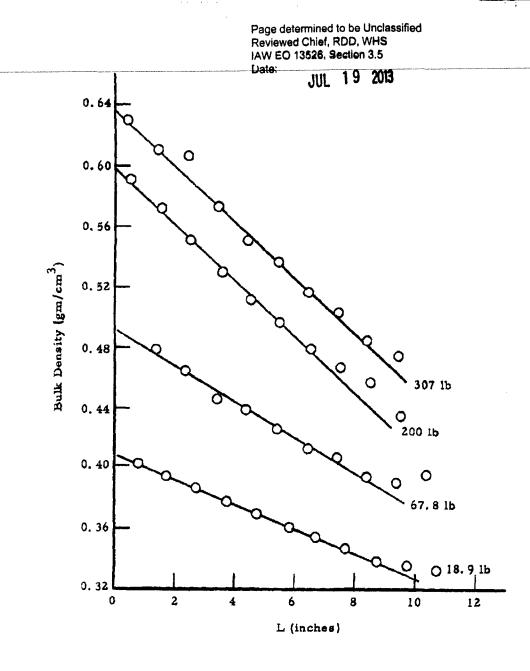
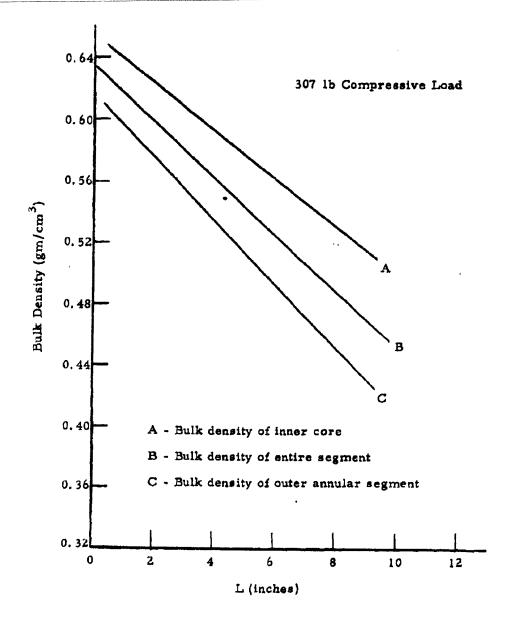


Figure 2.23 Bulk Density of Virgin Saccharin as a Function of Distance from Piston at Various Compressive Loads

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Figure 2. 24 Variation in Bulk Density of Compacted Saccharin along the Radius of the Plug

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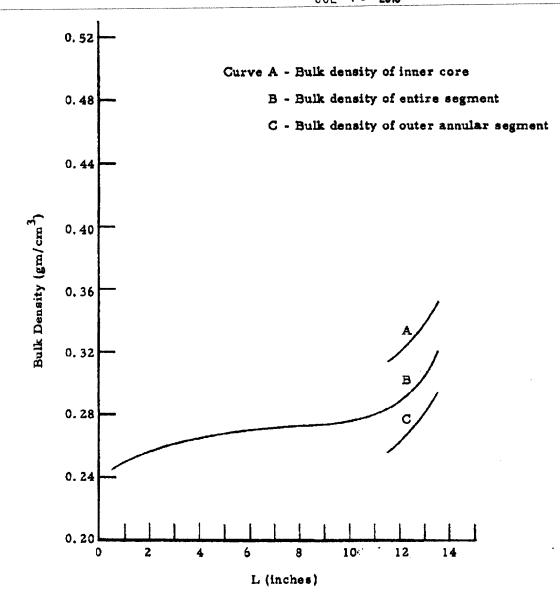


Figure 2.25 Variation in Bulk Density of Uncompacted Saccharin along the Radius of the Plug

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Further examination of Figure 2. 25 shows a sharp rise in bulk density near the bottom of the column of uncompacted powder. The deviation from linearity in the plots shown in Figure 2. 23 could be explained on this basis.

In attempting to explain the increased bulk density near the center of the compressed plug, one might suspect that during the filling process the powder tends to "pile up" in the center. Visual observations made during the filling process indicate that this is probably not the case. The phenomenon that is observed is a buildup of powder, apparently due to electrostatic charge on the particles, along the walls of the apparatus as shown in Figure 2.26. Not only could this lead to partial segregation according to particle size, but the powder along the walls is probably of lower bulk density since it is not supporting a mass of overlaying powder as is the material at the bottom of the cylinder. A sketch of this concept, involving a dense core surrounded by less dense material, is presented in Figure 2.26.

Future work will certainly include continued investigation in this area of proper packing of the column. We also plan to extend our studies to other powders such as tale, cornstarch, and powdered milk. There is, of course, a limitation in our selection of powders for study because of the relatively large volumes of powder required if virgin powders are used for each test.

## 2. 6 Shear Strength of Powders by Sliding Disk Method

During the present quarter our studies of the variation of shear strength of powders with compressive stress at various humidities have been extended to powdered sugar and powdered milk.

The techniques of measurement used were identical to those described in a previous report in which the powder, after exposure for at least 48 hours to a controlled humidity environment, is caused to shear while under the influence of a compressive stress by applying a shearing force normal to the compressive stress.

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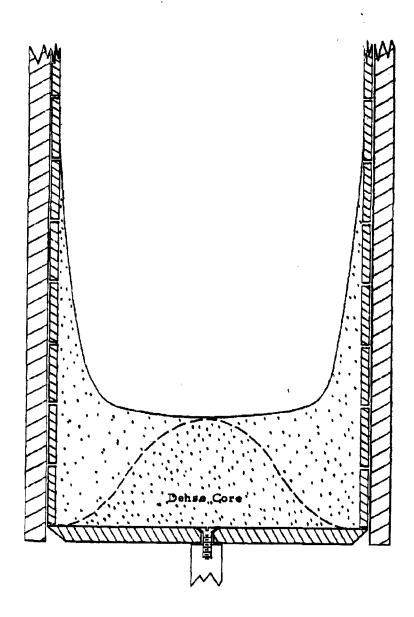


Figure 2.26 Sketch of a Probable Filling Mechanism in the Bulk Density Apparatus

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The results are tabulated in Figures 2.27 through 2.31. It should be noted from the graphs that the relationship between shear strength and compressive stress remains relatively unchanged for both powders during the increase in relative humidity from 2 to 32 percent. However at 46 percent relative humidity and more noticeably at 69 percent relative humidity there is a marked increase in shear strength with compressive stress. Also, the relationship between shear strength and compressive stress is no longer linear for powdered milk.

Future work will include studies of powdered sugar and powdered milk at higher compressive stresses. In particular it will be interesting to determine if the shear strength is in fact reaching a maximum value for powdered milk at 69 percent relative humidity as indicated by the plot in Figure 2.31. We also plan to extend our studies to other powders such as cornstarch, tale, and saccharin.

## 2.7 Future Area's for Study on the Characteristics of Powders

The emphasis in our work has been devoted to the development of reliable test methods based upon a sound theoretical background. In the future we will concentrate upon coordination of information on specific powders in order to completely characterize the powder, and to determine which test or tests yield the most reliable information concerning the compactibility and dispersability of a powder. New areas of study which will be initiated include:

## 2.7.1 Bulk Density of Loose Powders

An apparatus is being constructed in which powders will be allowed to fall a fixed distance through vibrating screens into a unit cube. It is believed that this will afford a simple, rapid, and accurate method for measurement of the bulk density of a variety of loose powders.

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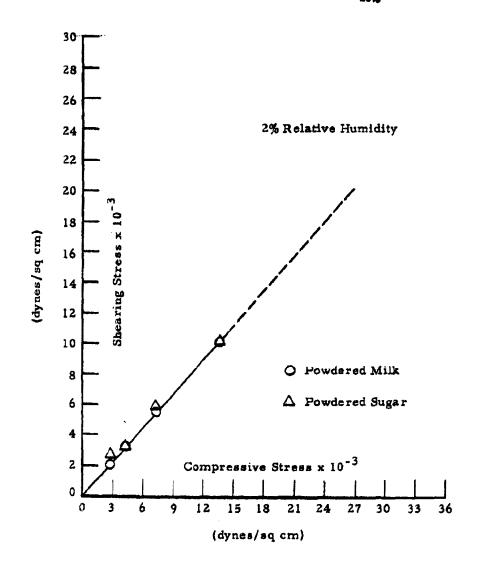


Figure 2.27 Variation of Shear Strength with Compressive Stress at 2 Percent Relative Humidity

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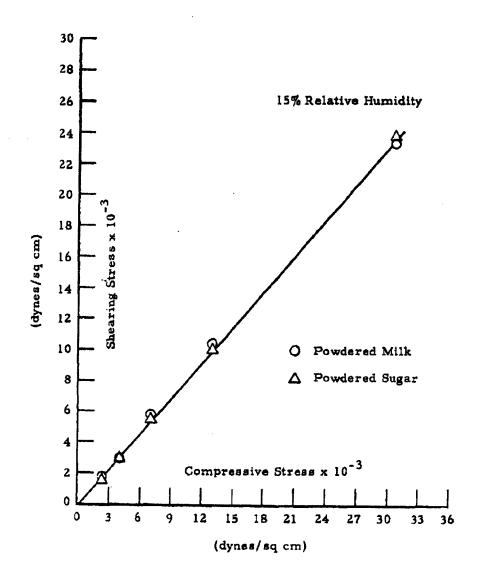


Figure 2.28 Variation of Shear Strength with Compressive Strees at 15 Percent Relative Humidity

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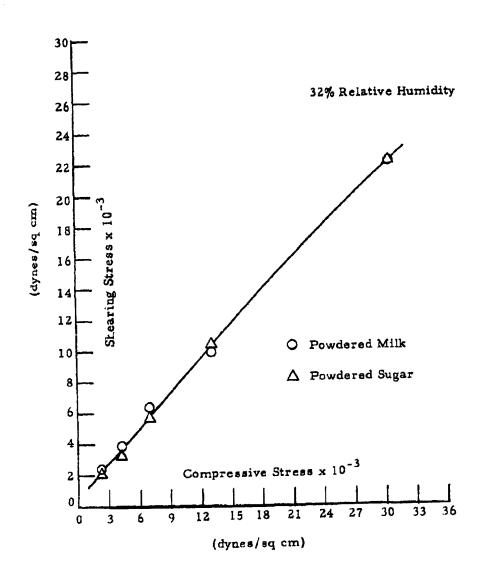


Figure 2. 29 Variation of Shear Strength with Compressive Stress at 32 Percent Relative Humidity

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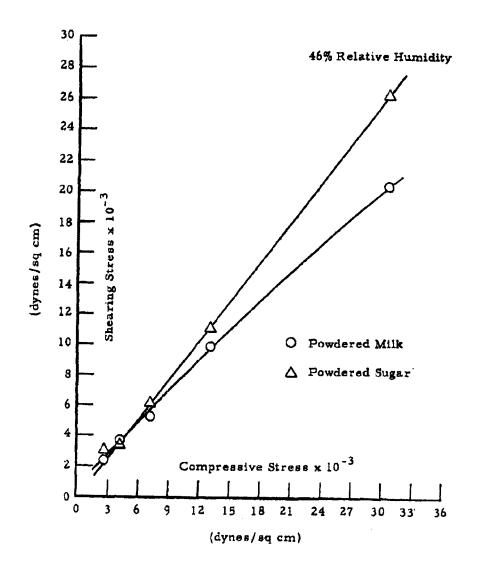


Figure 2.30 Variation of Shear Strength with Compressive Shear at 46 Percent Relative Humidity

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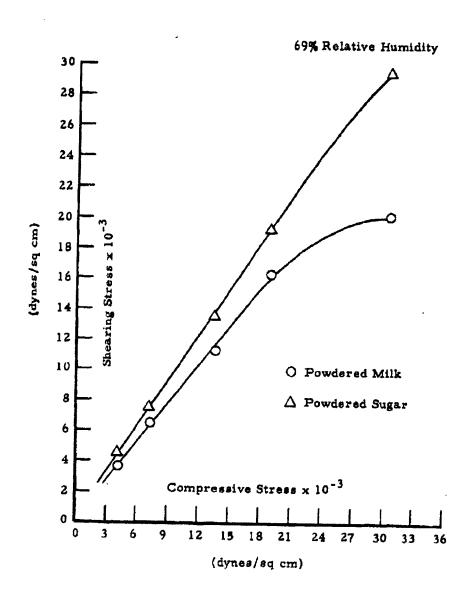


Figure 2.31 Variation of Shear Strength with Compressive Stress at 69 Percent Relative Humidity

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# 2.7.2 Measurement of Static Charge of Particulate Materials

A study will be made of various methods of generating and removing static charges, and of methods for qualitative and quantitative measure of static charge.

## 2.7.3 Particle Size and Shape

One of the most important characteristics determining the behavior of powders in the 5-micron range is the magnitude and the character of the total surface of the individual particle. A characteristic which can be called "surface roughness" should be of fundamental importance in predicting the ease with which a powder can be compacted and redispersed. This characteristic can be evaluated by a summation of information obtained from particle size distribution, particle shape, and total surface area. Particle size distributions are now being determined via the Whitby centrifuge. Photomicrographs are now being utilized to study particle shape. This study will be extended to the use of electron micrographs.

## 2.7.4 Total Surface Area

Measurement of total surface area of powders in conjunction with the particle size and shape analysis should yield a clearer picture of the total "surface roughness."

There are several methods for measuring the specific surface area of powders 6. Of these, the two most frequently used are the gas-adsorption (usually nitrogen) and air permeability methods.

In the gas adsorption method, the quantity of gas necessary to form a monomolecular layer on the surface is determined. By assuming a value for the area occupied by a single molecule, the area of surface covered by the

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adsorbed gas molecules is then calculated. The area measured by this method depends on the size of the molecules adsorbed and the degree to which they are able to penetrate into any cracks or pores in the solid.

The air permeability method is essentially a rheological method in which the flow of a fluid (liquid or vapor) through a powder in a compacted bed is related to the surface area of a continuous solid. It is assumed in this method that the bed of powder behaves as a bundle of capillaries. In consequence, only the surface of the continuous paths through the material will contribute to the measured specific surface area. This area is not the same as that measured by adsorbing a gas on the surface of the powder, because in the latter method all the surface accessible to gas molecules of the type used will contribute. In general, the area measured by the adsorption of a gas will be larger than that obtained by a permeability method. This difference may be accentuated if the powder has appreciable "internal" surface due to cracks, internal pores, and other irregularities. In addition, the permeability method may not measure the full "external" surface of the powder because of the formation of blind pores during compaction of the bed.

With very fine powders a further complication arises since it is very difficult to compact a fine powder to give a bed of low-voids fraction; and with a porous type of bed there may be a serious lack of uniformity throughout the compacted bed. From these considerations it would appear that the logical method to be used for powders currently under study would be the gas adsorption method.

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#### 3. AEROSOL STUDIES

The objective of this experimental program has been to perform a study of the properties of aerosols, both intrinsic and extrinsic, which have an effect on the aerosol's over-all stability. The degree of stability can be measured by observing the net rate of disappearance of the particles from the aerosol. Several factors, besides concentration, conceivably have an effect on aerosol stability. These factors are:

- 1) Partizle size and shape
- 2) Particle type
- 3) Charge characteristics of particles and environment
- 4) Water content of particles
- 5) Absolute humidity of environment
- 6) Type of settling (turbulent or tranquil)

The instability of aerosols is obviously to be understood in terms of only two distinct processes:

- 1) Particles may be removed from the aerosol by settling on and adhering to the various surfaces to which the aerosol is exposed.
- 2) Particles may be removed from the aerosol by colliding with and adhering to other particles within the same aerosol resulting in a net decrease of independent particles (coagulation).

Since the rates of these two processes completely specify the stability of an aerosol, the fruitfulness of any experimental method is to be judged primarily on its capability for supplying actual numerical values for these rates. Much of our work during this period has been directed at this type of evaluation of our light-instrumented aerosol chamber.

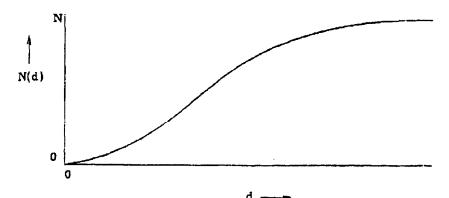
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#### 3. 1 Terminology and Definitions

A perusal of the literature of aerosols reveals a certain degree of independence among the various authors in regard to terminology. It is necessary, therefore, to define those terms used in this section which would conceivably conflict with the literature.

The number of particles per unit volume will be termed the concentration of the aerosol. The term concentration will often be specialized to mean the number of particles of a particular species, e.g., those with dimension in the range (d, d + od) per unit aerosol volume.

The classification of aerosol particles according to species warrants some comment. Since the most important particle property with respect to its behavior in an aerosol is particle size, the primary classification should be made accordingly. In dealing with solid particles, however, it is often not obvious how this classification should be put into practice because the particles are often very irregularly shaped. In our work we shall generally suppose that the particles may be considered spherical and characterize particle size by means of the diameter d. Thus if N denotes the total number of particles under consideration, we denote the number of particles with diameter less than d by N(d). Clearly N(0) = 0 and N(d) approaches N as d becomes infinite, so that a plot of N(d) against d is qualitatively as shown in the sketch below.



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The value  $d_{m}$  of d for which  $N(d_{m}) = 1/2 N$  is, of course, the median diameter. The terms monodisperse and polydisperse are widely used in reference to the variety of particle sizes present in a sample, a practice that we shall follow by saying that the dispersity is measured by the slope of the N(d) curve.

It is often convenient to speak of the fractional number of particles in a particular sample which have diameters less than d. Denoting this fraction by n(d) we have as its defining relation Nn(d) = N(d).

The derivative of N(d) has significance. The number of particles having diameters in the small interval (d, d +  $\delta$ d) is N(d +  $\delta$ d) - N(d) =  $\frac{N(d)}{\delta d} = N'(d) \delta d$ . Here the prime denotes the derivative. n'(d) is similarly defined. N'(d) is commonly called the particle size distribution while N(d) is often called the cumulative particle size distribution.

There are several explicit formulae for the particle size distribution, each of which approximates a wide variety of distributions actually encountered in practice. Each of these formulae contain two or more parameters which are supposedly capable of accounting for practical variations. Inasmuch as it is often convenient to have such a formula available for discussion, we have chosen one for that purpose. Since it is felt that the data in most cases are not precise enough to form a basis for choice of formula, we simply take the most common such formula which is compatible with the conditions n(0) and  $n(\infty) = 1$ , namely, the logarithmic-normal distribution:

$$n(d) = erf(\frac{\log (d/d_g)}{\log \sigma_g})$$

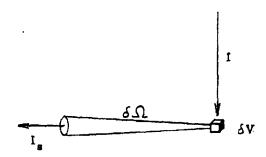
$$n'(d) \delta d = \frac{1}{(2 \pi)^{1/2} \log \sigma_g} \left\{ -1/2 \left( \frac{\log (d/d_g)}{\log \sigma_g} \right)^2 \right\} \delta(\log d)$$

Simple considerations show that  $d_g$  (the geometric mean diameter) in the above formula is also the median diameter mentioned earlier. The term  $\sigma_g$ , (the geometric standard deviation) is a measure of the dispersity.

#### 3.2 A Theoretical Calculation for Non-Agglomerative Aerosol Decay

It is expected that the information provided by scattered light will be rather subtly related to the fundamental processes of aerosol decay. For this reason it will be useful to have a simple, yet pertinent, relation between these processes and the aerosol light-scattering behavior. Such a relation is proposed in this section.

The light-scattering problem for an acrosol may be formulated as follows: Consider a small element of volume  $\delta V$  in the aerosol, this volume being irradiated from above by a monochromatic light beam of intensity I.



The amount of light scattered into a cone of solid angle  $\delta\Omega$  located at 90 degrees by a single particle of diameter d in  $\delta V$  is proportional to I and to  $\delta\Omega$ :

$$I_{\text{one particle}} = \sigma(d) \cdot I \cdot \delta \Omega$$

The constant of proportionality  $\sigma$  is called the differential scattering cross-section. Since the number of such particles in  $\delta V$  is  $C(d) \cdot \delta V$ , the total light scattered by particles of diameter d is:

$$\sigma(\mathbf{d}) \cdot \mathbf{C}(\mathbf{d}) \cdot \mathbf{I} \cdot \delta \mathbf{V} \cdot \delta \Omega$$

The light scattered by all particles is

$$I_g = I \cdot \delta V \cdot \int_{\sigma(d)} \sigma(d) \cdot C(d) \, \delta d \, \delta \Omega$$

This integral is to be evaluated under various conditions.

Let us first consider C(d). As has been explained in Section 3.1, the particle size distribution will be assumed logarithmic - normal. We further assume that at some initial time t=0 the concentration is uniform throughout the chamber. Thus:

$$C(d, t = 0) = \frac{N^{t}(d)}{V}$$

where V is the volume of the chamber.

This initial concentration may be modified with passing time. We consider two idealized modes in which the modification may come about.

First, suppose that the atmosphere in the chamber is "tranquil", i.e., sufficiently free of currents so that all particles settle vertically downward. In this case all particles of diameter d will fall as a unit, with the result that after time t, no particles of size d (or larger) will be present above a certain horizontal plane. This plane lies a distance h from the top of the chamber, h being given by:

$$h = t \cdot v(d) = t \cdot \frac{\rho_{gd}^2}{18\eta}$$

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where v(d) is the Stokes' settling velocity of a particle of diameter d,  $\rho$  is the particle density, g is the gravitational constant, and  $\eta$  is the air viscosity. The concentration as a function of time is therefore

$$C(d, h, t) = \begin{cases} C(d, t = 0) \text{ for } h > t \cdot v(d) \\ 0 \text{ for } h < t \cdot v(d) \end{cases}$$

that is, C(d, t = 0) is to be multiplied by a function which has value 1 for  $0 < d < \sqrt{\frac{18 hh}{\rho gt}}$  and value zero for larger d.

As an alternate mode of decay, suppose that the air in the chamber is so turbulent that the particles move in an essentially random manner. For this case it has been stated by various authors 7,8 that concentration remains uniform throughout the chamber, but that after time t the number of particles of size d is reduced by the factor

$$e^{-\frac{v(d)}{H}t}$$

where H is the total height of the chamber. Thus

$$C(d,t) = C(d,0) \cdot e^{-\frac{V(d)}{H}t}$$

The remaining factor in the scattering integral, the differential scattering cross-section  $\mathcal{O}(d)$ , is in general a very complicated function of d as well as of the light wavelength and the index of refraction. However, the ratio of particle diameter to light wavelength to be encountered in the present experiment is probably large enough so that a simple approximation from classical optics may suffice  $^9$ . This approximation is:

$$\sigma(d) = \frac{d^2}{6\pi}$$

The light-scattering problem thus leads to the two integrals: For the tranquil case:

$$\frac{1}{6\pi} \cdot \frac{\text{NI } \delta \text{V } \delta \Omega}{\text{V } \sqrt{2\pi} \ln \sigma_{\overline{g}}} \int_{0}^{\text{d max}} d^{2} e^{-\frac{1}{2} \left(\frac{\ln d/dg}{\ln \sigma_{\overline{g}}}\right)^{2}} \delta(\ln d)$$

with 
$$d_{max} = \sqrt{\frac{18 \text{ hm}}{\rho \text{gt}}}$$
;

For the turbulent case:

$$\frac{\text{NI} \delta \Omega \delta V}{6 \, \text{TV}} \cdot \frac{1}{\sqrt{2 \pi} \ln \sigma_g} \int_{0}^{\infty} d^2 e^{-\frac{V(d) t}{H}} e^{-\frac{1}{2} \left(\frac{\ln d/dg}{\ln \sigma_g}\right)^2} \delta(\ln d)$$

It is to be noted that the present considerations take no account of agglomeration. Another inadequacy, which may be partially corrected, is that the turbulent decay expression:

$$-\frac{v(d)}{H}t$$

takes into account only "gravitational impingement" on the floor and neglects "inertial impingement" on other surfaces of the chamber. The effect of aerosol decay by inertial impingement may be seen qualitatively by the following considerations.

As a current of air approaches a wall it is turned back, with an attendant centrifuging action. Thus, the particles experience an equivalent gravitational force, and inertial impingement is actually quite similar to gravitational impingement. The effect on the decay rate is to add an apparent gravitational constant g' to the g which appears in the exponent of  $\frac{v(d)}{H}$  t. For further details on this process, see Reference 8.

The integral for the tranquil settling case may be solved analytically. The result is:

$$d_{g}^{2} e^{2 \ln^{2} \sigma_{g}} \left\{ 1 - \operatorname{erf} \left( \frac{\ln \frac{v(d_{g}) \cdot t}{h} + 4 \ln^{2} \sigma_{g}}{2 \ln \sigma_{g}} \right) \right\}$$

$$= d_{g}^{2} e^{2 \ln^{2} \sigma_{g}} \left\{ 1 - \operatorname{erf} \left( \frac{\ln \frac{v(d_{g}) t}{h}}{2 \ln \sigma_{g}} \right) \right\}$$

where:

$$d_s = d_g e^{2 \ln^2 G_g}$$

is the surface median diameter. This function is linear if plotted in logarithmic-normal form with log t as abscissa. The 84, 50 and 16 percent points come at

$$t = \frac{h}{v(d_g) \sigma g^2}, \frac{h}{v(d_g)}, \frac{h \sigma_g^2}{v(d_g)}$$

respectively.

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The turbulent settling integral may be considered qualitatively by comparison to the tranquil expression. The two integrals differ in the modulating factor in the integrand. Where the tranquil integrand has a factor which drops from 1 to 0 sharply at  $\sqrt{d} = \frac{18 \text{ h} \text{ m}}{\text{Og t}}$ , the turbulent integrand

i di

has a factor (e ) which drops from 1 to 0 gradually, reaching 60.6 percent (the inflection point) at:

$$\frac{v(d)}{H}t = \frac{1}{2}; \quad d = \sqrt{\frac{18 \, \text{ft}}{\rho \, \text{g}^{\, \text{t}}} \cdot \frac{H}{2}} \quad . \label{eq:vd}$$

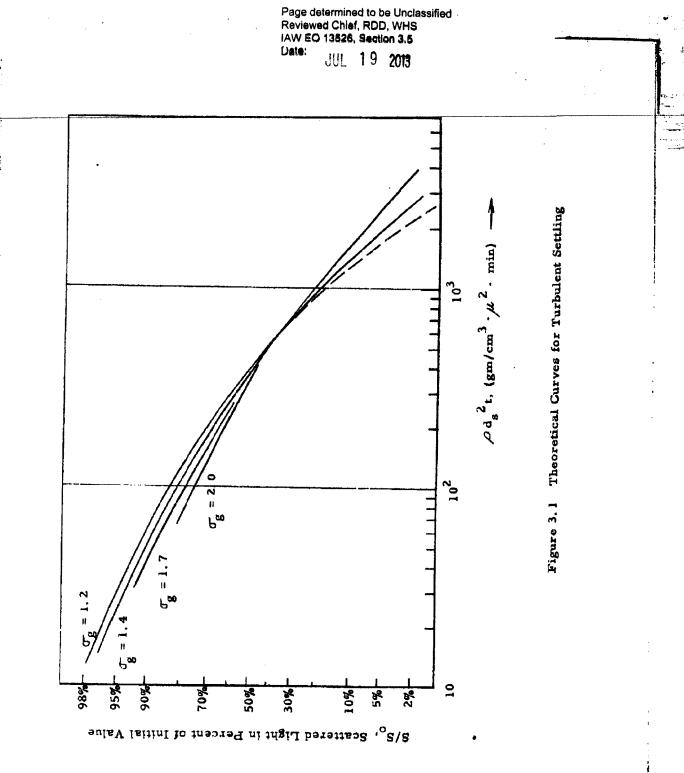
Thus if N'(d) changes sufficiently slowly with d (the aerosol is sufficiently polydisperse) the turbulent settling curve should be very similar to the tranquil expression if, in the latter, h is taken equal to  $\frac{H}{Z}$ .

As a check on the reasoning of the above paragraph, the turbulent settling integral was submitted to the computer group for numerical integration. A few curves, thus obtained, are shown in Figure 3.1.  $\rho d_s^2 t$ , with  $\rho$  in gm/cm<sup>3</sup>,  $d_s$  in microns and t in minutes, is used as the variable. It may be seen that the plots have considerable curvature for small values of  $\sigma_s$ , but approach linearity for larger values. The dashed line in Figure 3.1 is the plot of

$$-\frac{v(d_s)}{H} \cdot t$$

which is the solution of the turbulent settling integral as  $\sigma_g \rightarrow 1$ . It will be noted that latter nearly coincides with the curve for  $\sigma_g = 1.2$ . All the curves of Figure 3.1 cross 50 percent in the neighborhood of  $\rho d_g^2 t = 370$ , which may be compared with the value 286 obtained from

$$v(d_s) \cdot t = h \propto \frac{H}{2} \text{ at } 50\%$$



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#### 3.3 Experimental Work

The experimental objectives, somewhat idealized, of this program were stated earlier. As mentioned before, most of the work of the present quarter was carried out with a view to seeing how nearly these objectives could be accomplished in practice. Other areas of work were experimental techniques and determination of operating parameters.

Originally, it was considered desirable to restrict the scope of the experiment to the stability properties of aerosols; that is, to the behavior of aerosols subsequent to formation. This, however, requires the existence of a well-defined initial condition of the aerosol, a situation which is difficult to achieve in practice. Thus, while most of the runs to be discussed are concerned with aerosol decay (for which runs the dispersing process was assumed to be reasonably efficient), some work was directed at evaluation of dispersing efficiency.

The powder dispersing system was described in an earlier report. Briefly, it consists of a small chamber, containing the powder sample, which is pressurized with dry nitrogen gas. A diaphragm ruptures, due either to the pressure differential or to mechanical puncturing, suddenly releasing the pressure. A dispersing action is attendant. It was found that a 2-mil film of "phenoxy 8" plastic made a suitable diaphragm material, rupturing at 125 psi in our configuration. This material was used almost exclusively.

There were, broadly speaking, three series of runs:

- 1) Runs in which the fan was not used at all; these were intended for the study of the dispersing process.
- 2) Runs for which the fan was used only for a short initial period, intended for study of aerosol behavior under "tranquil" conditions.
- 3) Runs for which the fan was left on throughout the entire run, intended for study of zerosol behavior under "turbulent" conditions.

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It was found that stirring (Series (3) above) has a decided regularizing influence on the aerosol. In the "fan on" situation the two light-scattering signals were brought into coincidence almost instantaneously and stayed in coincidence, both decreasing smoothly, throughout the entire run. The Series (2) runs, on the other hand, often led to irregular (nonmonotonic) decreasing light signals subsequent to the turning off of the fan. The Series (3) runs will be discussed first.

Representative runs from Series (3) are shown plotted in logarithmic-normal form in Figures 3. 2, 3. 3 and 3. 4. Note that these plots are at least approximately linear. For purposes of discussion, we identify three abscissae, t', t\*, t", on this line (see Figure 3. 2). These are the intercepts of the line with 84, 50 and 15 percent scattered light intensity, respectively\*.

Table 3.1 presents a summary of the "fan on" runs. Note that t"/t', which according to Section 3.2 measures aerosol dispersity, is sensibly constant for a given powder while t\*, which is an inverse measure of the aerosol mean particle diameter, decreases with both amount of powder dispersed and with stirring. It may also be significant that the curves for the lower fan speeds differ more from linearity than the other figures. This point bears further checking.

The last two columns of the table show values of  $\rho d_s^2$  and  $\sigma_g$ , where  $\rho$  is particle density in gm/cm<sup>3</sup>,  $d_s$  is the surface median diameter in microns and  $\sigma_g$  is the geometric standard deviation, computed from t\* and t"/t'. The relations used are:

$$H = 2 \cdot v(d_g) t^*; \rho d_g^2 t^* = 286$$

<sup>\*</sup>According to the tranquil settling expression of Section 3.2, the ratio t''/t', which measures the slope, has the value  $(\sigma_g)^4$ . This should remain approximately true for turbulent settling. An approximate relation for t\*, as discussed in Section 3.2, is

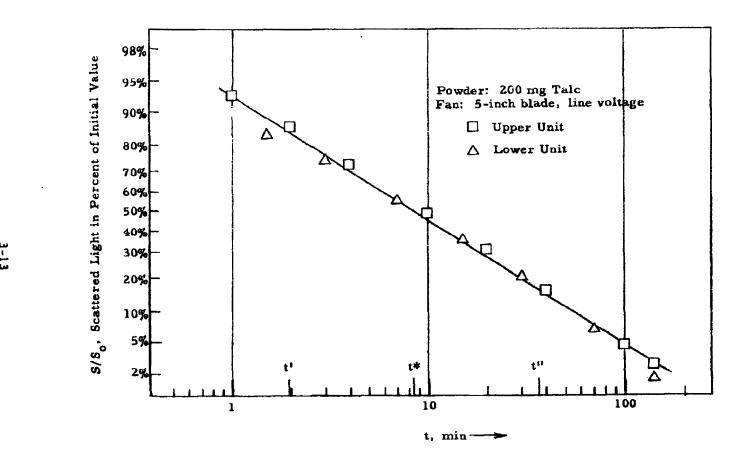


Figure 3.2 Experimental Curve for Light Scattered from Talc Aerosol

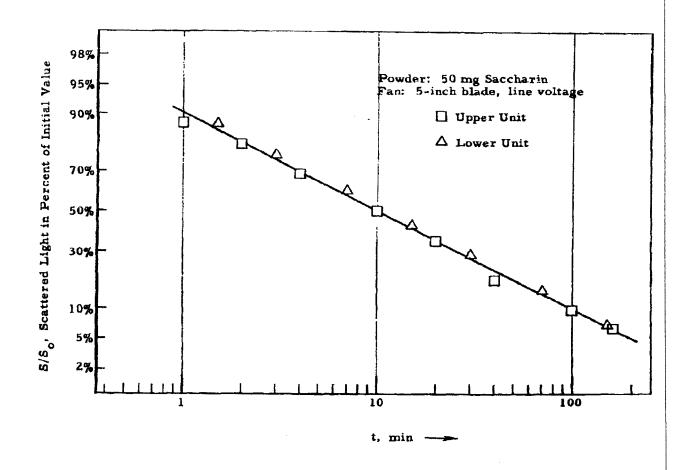


Figure 3, 3 Experimental Curve for Scattered Light from Saccharin Aerosol



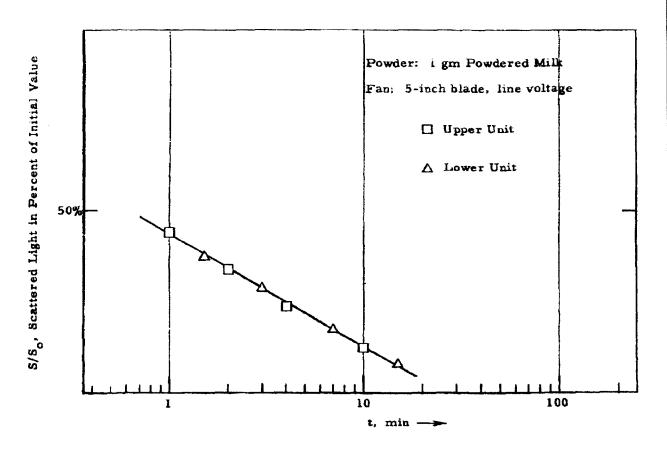


Figure 3.4 Experimental Curve for Scattered Light from Powdered Milk Aerosol

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$$\mathcal{O}\,\mathrm{d_s}^2 = \frac{370}{t^*}$$

$$\sigma_{\mathbf{g}} = \sqrt{t^{1/t^1}}$$

The values of  $\rho d_s^2$ , especially, may be viewed with suspicion since no effort has been made to take inertial impingement into quantitative account. Since our purpose is not aerosol assay as such, we do not intend to make the correction.

Table 3.1 Experimental Data for "Fan On" Runs

Powder Dispersed	Fan Condition	t* (min)	t''/t'	$\mathcal{O}_{d_g}^2$	σg
200 mg talc	Large blade, line voltage	8. 2	17. 5	45. 1	2.04
50 mg talc	Large blade, line voltage	12.2	21.5	30.3	2. 15
20 mg talc	Large blade. line voltage	19.0	20. 7	19.5	2. 13
50 mg talc	Large blade, line voltage	10.0	35, 3	37.0	2. 44
l gram pwd. milk	Large blade, line voltage	0:59	28.6	62.7	2. 31
56 mg talc	Small blade, 50 volts	20.5	20.4	18.1	2. 13
56 mg talc	Small blade, 35 volts	18.0	32. 8	20.5	2. 39

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Other "fan on" runs, conducted with yet a different powder, turned up an appreciably different type of light-scattering signal. The powder used here was "K ferric oxide", a very nearly monodispersed material of reasonably spherical particles. Particle diameter is approximately 2 microns. For this powder the scattered light decreased exponentially, as may be seen in Figure 3.5. This behavior is in accordance with the hypothesis of Section 3.2, Further, the experimental decay constant 0.035 compares favorably with the expected value of

$$\frac{v(d_g)}{H} = 0.037.$$

With regard to Figure 3.5, however, a question may be raised concerning the larger value of the slope during the first 5 or 10 minutes of each run. It is believed that there are several possible explanations for this rather minor deviation, none of which contradict the hypothesis advanced in Section 3.2.

Interpretive remarks, in terms of the relative roles played by agglomeration and by settling on surfaces, must remain rather hypothetical at this time. It has been stated that the light-scattering data for polydisperse powders are fairly well represented by a straight line on a logarithmic-normal plot. This would say that either the aerosols observed so far have log-normal particle distributions and decay without agglomeration (as in the model of Section 3. 2), or that agglomerative effects balance out the deviations from log-normality. It is also conceivable that the agglomeration has the simple effect of tilting or displacing the line on the log-normal plot, a behavior which was indeed observed. On the other hand, the line position

<sup>\*</sup>K ferric oxide was at one time produced and marketed as a pigment under the trade name "KROX" by Minnesota Mining and Manufacturing Co., St. Paul, Minnesota.

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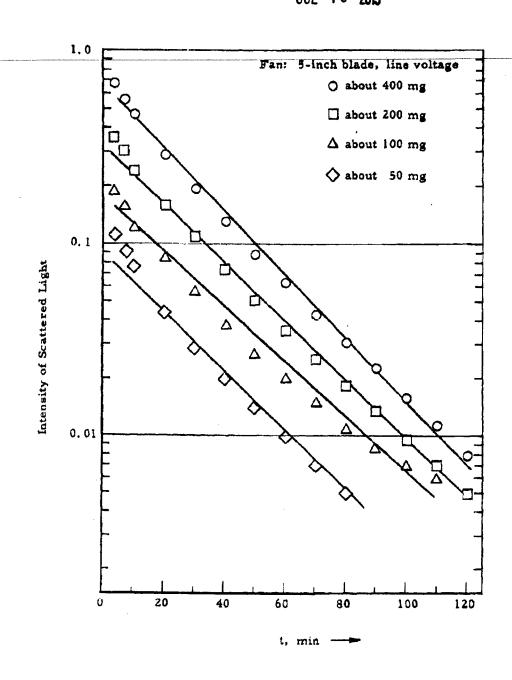


Figure 3.5 Experimental Curves for Light Scattered from K-Ferric Oxide Aerosol

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and slope may also depend on the efficiency of the dispersing process. All these possibilities will have to be studied further before more definite statements as to the relative roles of agglomeration and settling may be made.

Several runs have been made under "tranquil" conditions (Series (2) above). In view of the sometimes very erratic behavior of these aerosols, it would be misleading to list quantitative results; rather, some significant qualitative statements may be advanced.

The "tranquil" decay runs were made with the fan in operation from several minutes before to a few minutes (usually 3 or 6) after the firing of the powder charge. In general, the behavior was that the two light signals were equal and decreased smoothly until the fan was shut off. At this point the rate of decrease of scattered light changed markedly to smaller values and the two signals became unequal (the lower usually, but not always, exceeding the upper), both decreasing in a nonmonotonic way. Signal excursions of 20 to 30 percent above and below the median line, lasting several minutes, were common.

The implication of the erratic behavior is clear: "clouds" form in the aerosol and drift at random through the light-scattering area. The significant point is that these clouds form spontaneously, since the light signals decrease smoothly during the time the fan is on, indicating a uniform concentration throughout the chamber. These effects were particularly severe for the larger powder charges (above 100 mg). There may be a "threshold concentration", above which an aerosol is unstable and tends to form clouds.

One may speculate as to the mechanism of formation of the clouds. Perhaps in the absence of stirring, soft eddy currents, which tend to trap aerosol particles, are set up by convection effects. Another possibility is that a highly charged particle attracts others which, however, do not actually collapse on this nucleus to form an agglomerate.

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Another significant result of the "tranquil" runs is the <u>lack</u> of separation of the two light signals. According to Section 3.2 the lower signal should lag the upper by a factor of two in time. This was not observed, although in most cases the aerosols "tried" to follow the rule for the first half hour or so. Evidently there exist convection currents in the chamber which impose a turbulent condition on the aerosol, at least late in the life of the aerosol when the larger particles have settled out.

Finally, some remarks on the dispersing system are in order. A general feature of all runs to date has been the presence of a "spike" on the light-scattering record at the instant of firing. These spikes, which varied from 2 to 4 times the equilibrium value at t=0, were at first considered unimportant, arising perhaps from the mechanical shock of firing. During the course of the work, however, the speculation arose that spikes were due to a dense cloud passing the light-scattering region at high velocity. These high velocities, if present, are undesirable since it is likely that the powder is carried across the chamber and impacted on the floor opposite the dispersing gun.

Several runs were made without the fan (Series (1) above), incorporating various nozzles and other devices in an effort to slow down the dispersing gas stream. These runs generally tended to confirm the suspicion of a high velocity cloud since the relative amplitudes of the signals indicated that the aerosol was at first concentrated near the bottom of the chamber and "billowed up" from there during the course of the next half hour. None of the measures tried gave a really satisfactory improvement of the situation.

<sup>\*</sup>This behavior has been noticed in connection with other instruments - see Final Report, Vol. 4, Material Evaluation, GMI Contract No. DA-18-064-CM-2336, p. 26.

#### 3.4 Conclusions and Plans for Immediate Future Work

The work accomplished thus far has served to establish operating conditions and to test our understanding of the aerosol decay process. In regard to the latter, it has been seen that the theoretical expressions set down in Section 3.2 have a considerable degree of validity, and they have been found very useful in interpreting the light-scattering data. Their usefulness would, however, be enhanced if the effect of agglomeration in the aerosol decay were known. Efforts will be made to take account, at least qualitatively, of agglomeration.

The work of the past quarter has also turned up some unexpected possibilities. For example the "jitter" or "noise" on the fan-on traces appears to depend on the type of powder dispersed and in any case seems too large to be accounted for by electronic noise. Thus, the possibility arises that statistical fluctuations in the number of particles involved in scattering are being observed. It is conceivable that statistical methods may be applied to extract certain information from the "jitter" values, namely, actual numbers of particles involved in scattering. This possibility will be explored during the next quarter.

The experimental program of the next quarter will also be concerned with systematic studies of humidity effects in aerosol decay. As a first step several runs are planned under the two extreme humidity conditions, both of which should be relatively easy to establish. In these runs, as in others planned, we shall generally use stirring throughout the run, with the fan operating at speed sufficient to overcome convection currents.

The powder dispersing system will be kept under review. While this system does not work as planned, it is probably satisfactory for dispersing small charges of powder.

<sup>\*</sup>The same procedure was adopted by Tanner et al., Biological Laboratories Interim Report 36, "Design, Construction and Operation of the Aerophilometer" Special Operations Division, Camp Detrick, June 1953, p. 6.

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#### 4. VIABILITY STUDIES

#### 4. 1 Presence of Bg Contaminants in Sm Powder

The results of several trials designed to measure the degree of heat inactivation of Sm powder aerosols were invariably heavily contaminated with Bg colonies. Careful examination of our equipment and techniques eliminated these sources as potential causes. Hypothetical reasoning led to the conclusion that the Sm powder itself contained Bg contaminants in the ratio of  $1:10^6$  to  $1:10^7$ . This level of contamination would not be evident during routine platings of the Sm powder which had a viable count of  $1 \times 10^{11}$ /gm. When dilutions were made of this powder to yield countable plates (30 to 300 colonies), the plates would show only pure cultures of Sm. On the other hand, analyses of samples in which 99.999% of the vegetative Sm was destroyed by heat would yield plates in which Bg predominated.

This hypothesis was verified by preparing a liquid suspension of Sm powder in tryptose-phosphate broth which was immersed in a water bath at 75°C. Aliquots were removed at 5-minute intervals and plated in the appropriate dilutions to yield countable plates. The results showed a steady diminution in Sm counts relative to exposure intensity, and a constant level of Bg "contamination". At the end of 30 minutes the predominant organism on the plate was Bg.

This observation is significant only when considering circumstances where a given treatment would have different influences on vegetative cells compared to spores.

#### 4. 2 Effect of Agitation and Compaction on Viability of 5m Powder

A series of trials were performed to determine the effect of compaction on <u>Sm</u> powder viability, with and without the addition of Cab-O-Sil.

The additive was incorporated into the powder either by mechanical stirring

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or in a fluid energy mill. Immediately after compaction, plate counts were run both on the original bulk samples and on pellets disintegrated in a tissue homogenizer. These results are shown in Table 4.1.

Table 4.1 Effect of Agitation and Compaction upon Viable Count of Sm Powders

		Viable Count/gram x 1010		
Agitation	Additive	Bulk Powder	Compacted Pellet	Pellet Density
Untreated	none	4, 11	2. 84	0.617
Mechanical Stirring	none	3, 01	2. 15	0.624
	1% Cab-O-Sil	2, 72	2.69	0.629
Fluid Energy Mill	none	3. 13	3.39	0.625
	1% Cab-O-Sil	3, 07	2.68	0.616

It is evident that neither coating with Cab-O-Sil nor compaction to 0.62 grams per cc density had any significant or deleterious immediate effects on <u>Sm</u> viability.



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### 5. DISSEMINATION AND DEAGGLOMERATION STUDIES

#### 5.1 General

During this period major emphasis was placed on wind tunnel experiments at Fort Detrick employing the 40-foot diameter test sphere. The objective of this program was to evaluate the General Mills GMI-3 fixture with Sm and P. tularensis, under environmental operational conditions which simulate actual operation of the prototype unit for dissemination of dry BW agents. In the evaluation such parameters as recovery factor, source strength, decay rate, VMMD, slopes with 95 percent confidence limits, LD<sub>50</sub>, probit slope and its 95 percent confidence limits will be determined. The Sm test series was conducted on April 4, 5, 6 and 9, 1962, while the P. tularensis series was conducted on May 4, 7, 8, 9, 10 and 11, 1962\*.

In preparation for the above activity, effort was also devoted during the period to developing the test fixture in our laboratory.

# 5. 2 Evaluation of the General Mills, Inc. Test Fixture Disseminating Dry Sm and Dried Pasteurella Tularensis

At the present time statistical analyses of the results of the Fort Detrick test program are not available. However, the procedure and general results will be given in this report.

The over-all program was designed to test six operational treatments with dried Sm and dried P. tularensis. These included the two wind tunnel Mach numbers, 0.5 and 0.8, and three bulk densities: normal, approximately 0.43 gramsper cc and 0.53 grams per cc. In the Sm test four

<sup>\*</sup>Technical Evaluation Division Test No. 62-TE-1602; MD Division No. 1927.

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replicates were obtained per treatment, while in the <u>P. tularensis</u> series six replicates were determined per treatment. In all of the tests, each treatment was run daily.

Apparatus for these tests included, in addition to the GMI-3 fixture, related equipment used for compacting powders to the required bulk densities and mechanically breaking down the resulting slugs into fine agglomerates on the order of 2000 microns and less and basic particles. Approximately 0.8 grams of these samples were placed in the GMI-3 fixture and disseminated at a feed rate of 30 lb/min. These apparatus are further discussed in our previous report 10.

A blowdown wind tunnel and auxiliary equipment such as air storage tanks and pumps were erected for these experiments adjacent to the test sphere. The test section into which the organisms were disseminated, shown in Figure 5.1, has a 2-inch by 4-1/2-inch cross-section. Actual ejection of the material into the air stream was achieved during a four-millisecond period.

Environmental conditions inside the sphere during these tests were maintained at 75 degrees Fahrenheit temperature, essentially atmospheric pressure and 35 percent relative humidity.

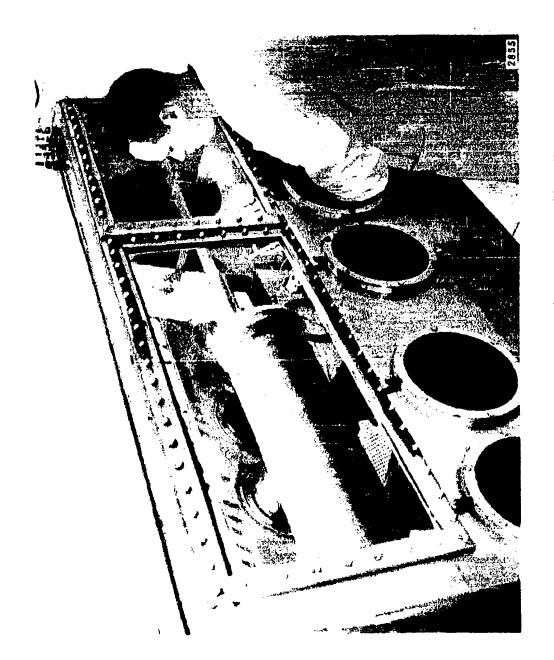
Samples of the generated aerosols were obtained with ABP-30 samplers\* filled with 20 milliliters of tryptose saline during Sm trials and 20 milliliters of gelatine peptone phosphate during P. turlarensis runs. One minute samples were taken at the midpoints of 4, 18, and 32-minute time periods at a flow rate of 12.5 liters per minute. Four sampling stations were operated concurrently.

Single stage impactors (SSIs) with cutoffs of 3, 5, 7, and 9 microns and a GP-20 total collector were used with 25 milliliters of sampling fluid. This group of samplers were operated only at the 4-minute time period.

<sup>\*</sup>All-glass impinger preceded by a British pre-impinger.

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Wind Tunnel Test Section Located in Glove Box of Spherical Test Chamber at Fort Detrick Figure 5. 1

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The four ABP-30 samples taken during each time interval were pooled while the SSIs and GP-20s were assayed separately.

In the original P. turlarensis test program, it was planned that eight Hartley strain guinea pigs in the weight range. 250 to 370 grams, would be exposed at each of three time periods for each trial after the first day. However, because of the high recovery factors during the initial experiments, a new test plan for the animals was required. As a result they were only exposed on the final three days.

#### 5.3 Preliminary Results

#### 5. 3. 1 Sm Tests

In addition to determining the viability of generated aerosols in the sphere, control samples of the normal and compacted materials were taken and assayed. By this method data were obtained on the effects of two factors which enter into this evaluation program. 1) the compaction of the particulate material to densities up to 0.53 grams per cc. and 2) the aerodynamic breakup process during dissemination. Each factor can be analyzed separately or the two can be combined to demonstrate the over-all performance for any one treatment.

In studying the mean values of the control counts, there appears to be little, if any, effect on viability due to compaction of  $\underline{Sm}$ . Count values ranging from 123.98 x 10<sup>9</sup> to 127.35 x 10<sup>9</sup> organisms per gram were obtained at bulk densities 0.53 grams per cc and 0.43 grams per cc, respectively. At normal bulk density a value between these was obtained.

Mean values of the recovery factors at the 4-minute time period varied randomly in a range from 2.52 to 3.36 percent for the six treatments, while for the AP #10 fixture the value was 3.50 percent. At the two wind tunnel Mach numbers studied, there appears to be little difference in the viability of the generated aerosol.

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At first glance these recovery factors may be deceiving and therefore may appear quite low. Thus, it is necessary to explain that the recovery factor is directly proportional to the quantity of fill which has a particle size less than 5 microns. In this case the particle size distribution, based on a Whitby centrifuge analysis, showed that only 6.5 percent by mass of the material consisted of particles less than 5 microns in size. In addition, the mass median diameter and geometric standard deviation of the material was 16.6 microns and 1.95, respectively.

#### 5.3.2 P. Tularensis Tests

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Preliminary data similar to that presented above are also available for the trials with P. tularensis. In the case of the control samples, the mean values at each bulk density, based on five replicates, range from 13.52 x 10<sup>10</sup> to 20.11 x 10<sup>10</sup> organisms per gram for bulk densities ranging from the normal condition to 0.53 grams per cc. respectively. The results show an increase in organism count with increase in compaction, an effect which is unexplained at this time.

Mean recovery factors for this series of tests based on six replicates range from 28.59 to 35.77 percent. There is an indication that recovery decreases at the higher bulk density due to aerodynamic breakup; however, for the combined effects of compaction and breakup, increasing recovery factors were obtained with increasing bulk density. Again in this case, wind tunnel Mach number appears to have no relative effect on recovery.

The mean recovery factor obtained with the AP #10 reference fixture was 28.44 percent.

The mass median diameter of the sample material was 4.3 microns while the geometric standard deviation was 1.38. Approximately 70 percent by mass of the material was less than 5 microns in size.

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In spite of the fact that the particle size distribution was fairly fine, the recovery factor obtained in these tests is considered to be quite high and exceeded our expectations.

A more complete presentation of the results from both the Sm and P. tularensis tests will be made at a later date when the complete statistical analyses have been finished. At that time it should be possible to make definite conclusions as to the operational characteristics and effectiveness of the GMI-3 fixture and wind tunnel combination.

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## 6. CONTINUATION OF EXPERIMENTS WITH THE FULL-SCALE FEEDER FOR COMPACTED DRY AGENT SIMULANT MATERIALS

#### 6.1 Introduction

Experimental work with the full-scale feeder continued during this reporting period. The information obtained substantiates previously reported conclusions that the design concept under study is quite feasible. Performance was evaluated at powder flow rates ranging from 20 lb/min up to 49 lb/min. Air flow rates from three to nine sefm were investigated. Particular attention was devoted to performance during the starting period and various operating procedures were tried in an effort to minimize the delay in reaching full feeding rates.

In general, the procedures employed in conducting these test runs are the same as described in the Seventh Quarterly Progress Report <sup>12</sup>. A few modifications were made to the unit and an improved arrangement for collecting the discharged material was devised.

#### 6.2 Equipment Modifications

The original plan was to try three diameters for the discharge opening - 0.375, 0.500 and 0.750 inches. The smallest diameter of 0.375 inch was soon determined to be too small. The 0.500 and 0.750 diameters were both satisfactory but it was found that the gas pressure within the unit using the 0.750 diameter opening was significantly lower. Consequently, a decision was made to use the latter opening in all of the test runs being reported herein and to tentatively establish this diameter as the one to use in the airborne disseminator.

For the tests reported in the Seventh Quarterly Progress Report 13 the motivating gas was admitted directly into the disaggregator section of the unit through a hole near the top of the cylinder. For the tests reported herein, the unit was modified in that an aerating ring was installed in the

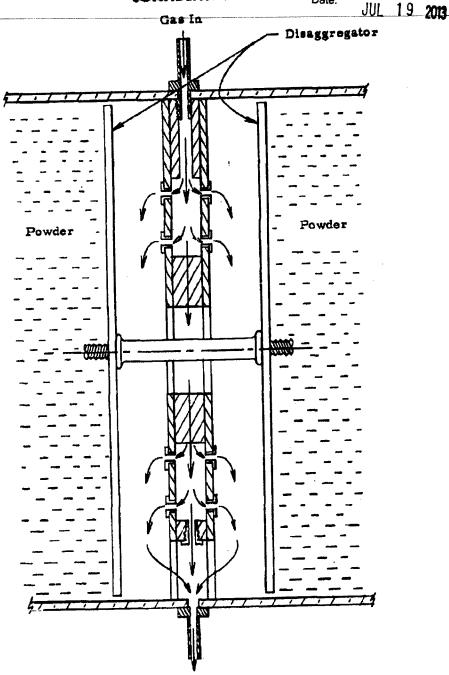
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center of the cylinder as shown in Figure 6.1. This ring contains 32 orifii each 0.040 inch in diameter which are positioned uniformly around the ring on both sides. Four additional crifii are mounted 1.75 inches directly above the powder discharge opening. The objective in using the aerating ring is to obtain better mixing of gas and powder. To accommodate this aerating ring it was necessary to remove the wire wall scrapers and the 6-inch baffles which were mounted between the disaggregator disks. The scrapers were eliminated but the four long baffles were replaced by eight 1.25 x 2-inch baffles, four being mounted on each disk.

The initial arrangement for collecting powder for weighing was to use a short length (approximately 24 inches) of 1.25-inch inside diameter tubing between the discharge on the unit and the top of the 50-gallon drum. The top of the drum was covered with a sheet of muslin to allow the gas to pass out of the drum without carrying out the powder. Since this technique could conceivably affect the powder flow at the point of discharge from the disseminator, a new technique was devised in which the tubing was emitted and the powder allowed to flow directly into a stand pipe six inches in diameter mounted on the collection drum as shown in Figure 6.2. The end of the stand pipe is placed four inches below the discharge tube on the disseminator. The cover of the drum is well ventilated by many holes and the muslin filter cloth is placed on top of this cover. A chamber is formed above the cover by means of plastic sheeting so that the gas can be withdrawn from between the plastic and the filter cloth using a vacuum cleaner. With this technique the powder has been observed to flow freely into the collection drum in a well-defined stream or jet.

#### 6.3 Test Results

The results obtained with the full-scale feeder during this reporting period are summarized below. This summary is followed by a more detailed discussion of the data.



Gas + Powder Out

Figure 6. 1 Aerating Ring Installed in Center Section of Full-Scale Experimental Feeder



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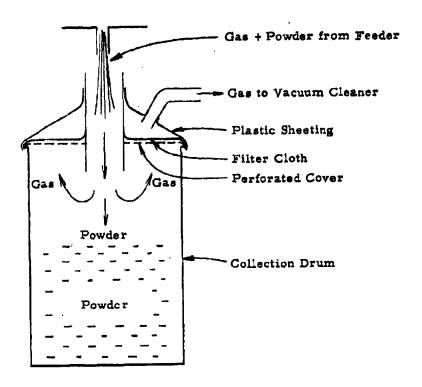


Figure 6.2 Arrangement for Collecting Powder without Direct Connection between Collection Drum and Discharge Tube



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A total quantity of approximately 2,600 pounds of talc were successfully loaded into and discharged from the feeder under a variety of operation conditions.

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- The torque and power required to drive the feeder were found to be well within the initial objectives. The torque was usually less than 100 ft-lbs and the power less than 0.5 hp.
- A discharge opening of 0.75 inch in diameter gives good 3) performance over the feed rates investigated.
- Powder was discharged successfully with gas flow rates ranging from three to nine scim.
- A gas flow rate of at least six scfm was found to be necessary in order to minimize the delay in attaining full powder discharge rate.
- The delay in attaining full powder discharge rate was also minimized by pre-pressurizing; i.e., the operating pressure was established in the cylinder before starting to feed
- The gas requirements were found to be compatible with the space which will be available for storing gas within the airborne disseminator.
- No significant difference in performance was observed for operation where the gas flow was started before feeding the powder as compared with operation where gas and powder were started simultaneously.
- The feeder operated satisfactorily for powder flow rates ranging from 20 to 491b/min.
- The gas pressure measured within the cylinder was found 10) to vary from approximately 0.3 to 1.1 psig depending upon the powder and gas flow rates used.

The experimental feeder was operated successfully under various conditions of powder and gas flow rates. The range of operating conditions covered is shown in Figure 6.3 by plotting each run as a point on a family of curves showing air-to-powder ratios (by weight) for various powder and gas flow rates. The highest powder feed rate obtained was 49 lb/min.

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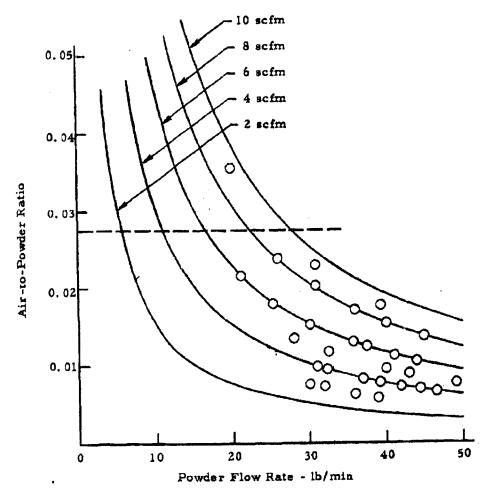


Figure 6.3 Air-to-Powder Ratios for Various Feeder Operating Conditions (actual test runs with talc are plotted)

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This is not the maximum capacity of the feeder but it is the highest rate attainable with the present variable speed drive when the feeder is loaded with material at a density of approximately 0.48 g/cc. Operation up to rates as high as 60 lb/min are planned by compacting to higher density during loading.

Typical powder flow rate curves are presented in Figure 6. 4. The plotted data were obtained by recording the time at which a certain amount of powder had accumulated in the collection drum. In most cases time was noted for each 5-lb increment. The curves were selected from many such curves which have been drawn to illustrate typical performance for a variety of operating conditions. The data for any given run produce a welldefined straight line indicating that the flow from the feeder is very uniform.

The driving torque and speed were observed for all test runs so that the input power requirement could be determined. The torque was observed to vary over a range from 15 to 100 ft-lb with one exception where the torque rose to 240 ft-lb. This excessive torque prompted an investigation which revealed that the thread clean-out feature which removes powder from the feed screw was plugged. This deficiency was corrected and there was no re-occurrence of an excessively high torque value.

The maximum driving speed used on any test was 24 rpm. The horsepower requirement based upon 24 rpm and 100 ft-lb is 0,46 hp.

One series of test runs was conducted using nitrogen rather than air as the motivating gas. As was expected, there was no observable difference in performance of the feeder when using nitrogen as compared to air.

The curves in Figure 6.3 can be used to show how the motivating gas flow employed in laboratory tests compares with the allowable flow when the gas supply is limited as it will be in the airborne disseminator. The design study has shown that it will be feasible to carry a gas storage bottle capable of supplying 113 standard cubic feet of nitrogen. When based on the maximum payload capacity of 350 lb, this gives a ratio of 2.75 percent by weight

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Material: "Mistron Vapor" Talc

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Compaction Density: 0.45 to 0.48 gm/cc

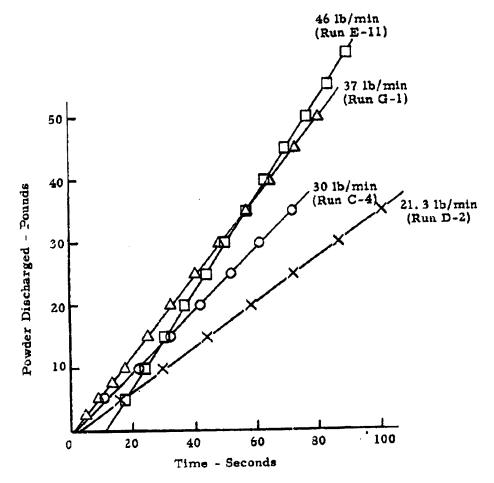


Figure 6.4 Powder Discharge Rate



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of nitrogen-to-agent payload. The horizontal dashed line in Figure 6.3 for this ratio is well above the conditions at which the feeder has been demonstrated to operate satisfactorily.

The gas is used most efficiently when the gas flow rate is reduced as the powder feed rate is decreased. However, it is possible to operate the feeder over a wide range of powder flow rates with a fixed gas flow rate without exceeding the 2.75 percent ratio. For example, if the gas flow were set at six scfm the powder feed could range from 17 lb/min up to 50 lb/min or more. In order to stay within allowable limits, the gas flow rate must be quite low when the powder feed rate reaches 10 lb/min. Anticipating a problem, the feeder performance was tested at the low gas flow rates.

In switching to the lower gas flow rates (three and four scfm) it was observed that there was a delay in reaching the full powder flow rate at which the unit was being operated. To obtain a measure of this delay, the flow rate curve is drawn as a straight line extended back to intersect the time-axis as shown in Figures 6.4 and 6.5 where time is measured from the instant at which the unit is started. It is desirable to obtain performance such that full flow rate is established in the shortest possible time.

Up to this point the operating procedure was to establish the desired gas flow before starting the drive system which feeds the powder. When operating in this manner the pressure within the feeder is essentially equal to atmospheric pressure before the powder starts feeding. When feeding starts the pressure increases to a value which depends upon both the powder flow rate and the gas flow rate. If the approximate operating pressure exists in the cylinder before starting the powder feeding, the delay in reaching full powder flow rate should be a minimum. Therefore, tests were made in which this internal pressure was established prior to feeding powder by restricting the air flow just below the discharge opening of the feeder. The restriction was removed at the same time that the powder feeding drive was turned on.

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Material: "Mistron Vapor" Talc

Compaction Density: 0.48 gm/cc

Cylindrical Pressure: 6.4 cm Hg

Air Flow: 4 scfm

Powder Flow: 43 lb/min

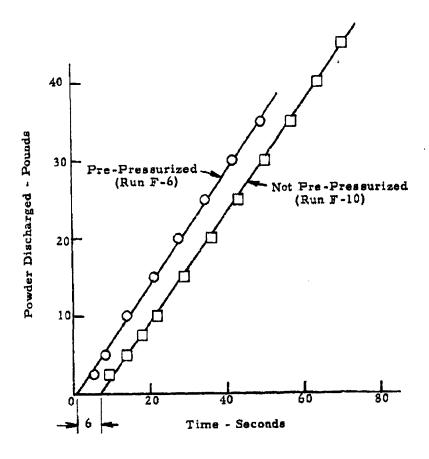


Figure 6.5 The Effect of Pre-Pressurizing on the Time Required to Establish Uniform Flow



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The pre-pressurizing technique proved effective as shown in Figure 6.5. These flow rate curves were obtained with four scfm air flow and 43 lb/min powder flow for both cases. The vertical separation of the two curves represents approximately five pounds of material. This indicates that without pre-pressurizing about five pounds more talc was stored in the disaggregator section of the feeder during the initial seven or eight seconds of operation for the test where pre-pressurizing was not used.

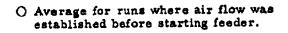
When operating the feeder for a series of runs in which the same powder feed rate was used for each run while the gas flow rate was varied, it was observed that the delay in attaining full powder flow rate, as determined from plotted flow rate curves, tends to be longer for the low gas flow rates than for the high. During these runs the following data were obtained:

- Time from starting feeder drive to the instant when powder began to flow from discharge opening.
- Time from stopping feeder drive to termination of powder flow.
- Amount of powder discharged after feeder drive was stopped.

The results of these observations are presented in Figures 6. 6 and 6.7 only to illustrate trends in performance characteristics; the amount of data involved is insufficient to justify using these curves to obtain lag-times or post powder-flow quantities for application in any related studies.

The time-lag data are presented in Figure 6.6. The lower curve shows how the observed time from start of feeder to start of powder flow was almost independent of the rate of gas flow used. The delay was approximately one second. The delay in stopping, however, increased significantly as the gas flow decreased below five scfm. The delay observed for eight scfm was approximately 2.25 seconds. It increased slightly to approximately 2.75 seconds at five scfm, but rose sharply to about eight seconds at three scfm.

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Average for runs where air flow was started when feeder was started.

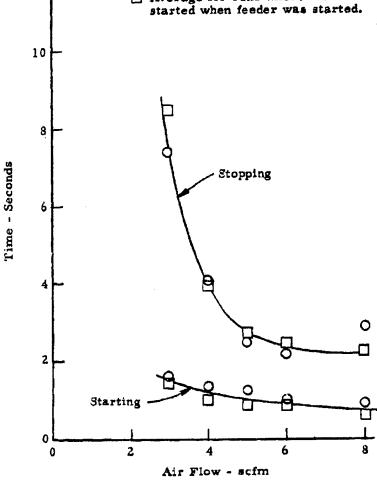


Figure 6.6 Time Lag in Powder Flow



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- O Average for runs where air flow was established before starting feeder
- Average for runs where air flow was started when feeder was started

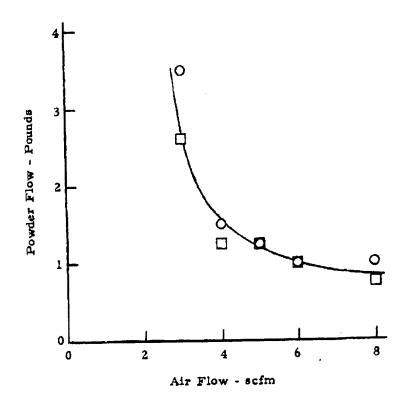


Figure 6.7 Powder Flow After Stopping Feeder Drive



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Figure 6.7 shows the amount of powder collected after stopping the feeder as measured for various gas flow rates. The curve shows that the air flow rate must be above five scfm if the amount of powder is to be minimized.

Another factor which could conceivably influence the initial delay is the relative timing of the start of gas and powder flow. This aspect of the problem was studied by making two types of runs, the first with the gas flow established prior to feeding powder, and the second with gas and powder flow started simultaneously. Data from both types of runs were used in plotting the curves in Figures 6, 6 and 6.7. The data show that there was no significant difference in performance for the two methods of operation. This was also true when performance was evaluated on the basis of flow rate curves drawn for the runs.

### 6.4 Plans for Future Work

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All of the experimental feeding experiments with the full-scale unit have been conducted with talc compacted to a density of approximately 0.45 g/cc. A hydraulic press is now being fabricated which will make it possible to compact to higher densities when filling the feeder. Tests will be performed at a density of 0.6 g/cc or higher when the press is available. This density will result in a powder flow rate of at least 60 lb/min when the feeder drive is set for maximum speed.

A supply of powdered sugar has been obtained and plans are to run the feeder with powdered sugar to determine if performance varies with the material being handled. Since powdered sugar is more hygroscopic, these tests will not be conducted until the air dryer for humidity control is installed and operating. The dryer installation has been delayed because the vendor failed to meet his quoted delivery date.



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Some of the future feeding experiments will be performed with the laboratory model of the disseminator which the Engineering Department is now fabricating. This unit is to be a full-scale prototype of the inner tank assembly of the airborne disseminator and will operate on the same basic principles employed in the unit which has been thus far used to demonstrate the feasibility of feeding compacted dry powder materials.



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# 7. Design studies on a dry agent disseminating store

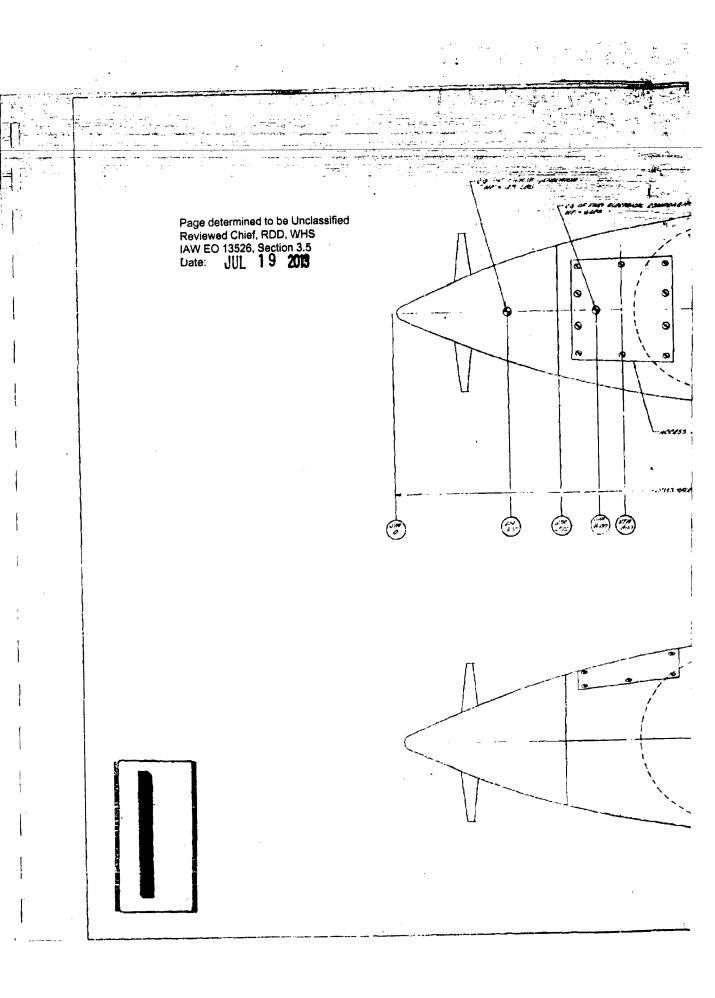
The design concept of an external aircraft store for disseminating dry agent material from a compacted state was described in our Fifth Quarterly Progress Report . Section 6 of this current report contains a discussion of the experimental evaluation of the basic concept using a full-scale model. This experimental program has demonstrated that the proposed concept is feasible. Some of the principal components of an airborne disseminator based on this concept were described in the Seventh Quarterly Progress Report. During this eighth reporting period progress was made toward integrating these components into a well-designed disseminating store as described in the following paragraphs.

# 7.1 General Arrangement for the Airborne Dry Agent Disseminating Store

A preliminary general arrangement for the airborne dry agent disseminating store is shown in Figure 7.1 (GMI Dwg SK-29100-612). The basic structure is an aluminum shell that provides a mounting skeleton for the inner tank assembly, the gas supply system, the turbine generator, the actuator and the necessary control apparatus.

The external skin is a 180-inch long aerodynamically shaped tank that has the same dimensions as defined for the 150-gallon Fletcher Aviation Co. Store No. 21-150-6024. This is a store that utilizes both 14- and 30-inch lug spacing for aircraft mounting. Two horizontal fins are mounted on the aft section to augment tank stability during flight.

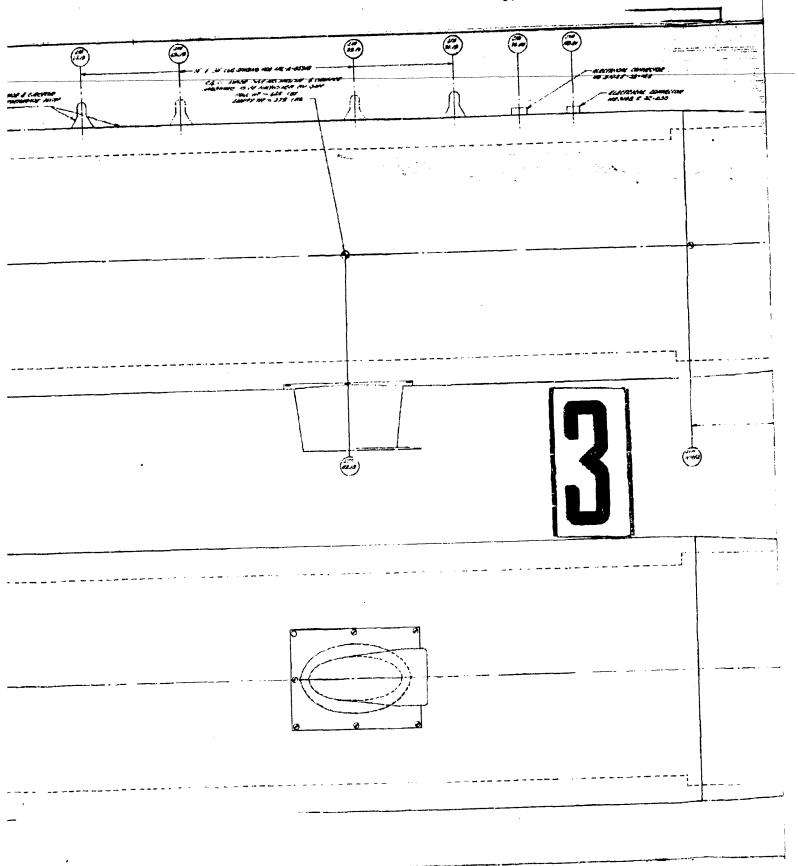
The store (exclusive of turbine generator assembly) consists of the forward, the center and the aft sections which extend from stations 13.0 to 52.0, stations 52.0 to 109.62, and stations 109.62 to 180, respectively. This particular sectioning of the tank was required to provide access to both ends of the inner tank for charging with the compacted dry agent. To minimize assembly time, the joints at stations 52.0 and 109.62 will be



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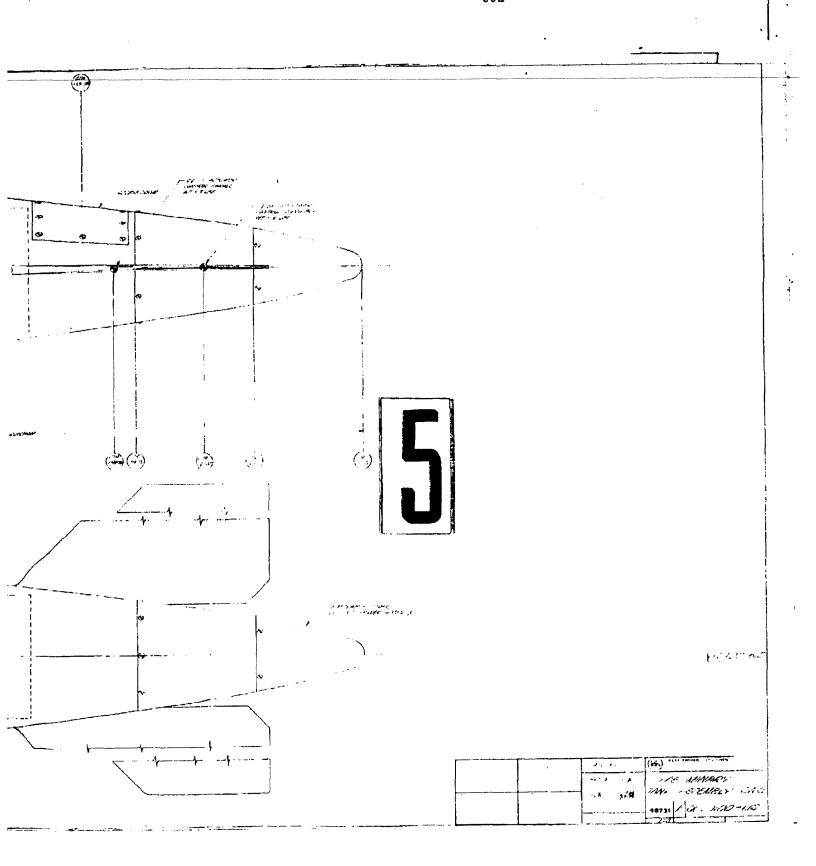
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equipped with a bayonet-type joining feature. In separating or joining the tank sections all that will be required is the loosening of a locking screw and rotating the end section approximately 22.5 degrees. Gas line and cable disconnects will be provided on the fore and aft sections from the center section. Access doors are located in the nose and tail sections at positions determined on the basis of necessity and ease of manipulation of the various system components and controls. Locations of various system components were evaluated during this period on the basis of available space, system operation and over-all effect on the store center of gravity.

The forward section from stations 13.0 to 52.0 contains the primary gas system hardware and the necessary electrical components for power distribution. The ram air turbins generator is mounted at station 13.0.

The center section, with section joints at stations 52.0 and 109.62, will provide the primary support structure and will house the inner tank assembly which contains the agent payload and the feeding mechanism.

The aft section from stations 109.62 to 180 contains the rotary actuator assembly, the actuator control electronics (including a control panel), and the ground and generator power distribution system electronics. Gas system control electronics are also included in this section. A special coupling is required between the actuator and the drive screw for easy separation when the aft section is removed for the dry agent loading operation.

#### 7.1.1 Ram Air Turbine Generator

The ram air turbine generator is the same 4.5 kvs, 115/200 volt, 400 cps, 3-phase generator made by General Motors. Allison Division, which is used on the GMI liquid agent disseminating store. However, it will be necessary to modify the exterior of the generator housing to make it conform to the shape of the dry agent disseminator. The generator is thoroughly described in GMI Specification GMS-29100-020, a copy of which was appended



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to our wifth Quarterly Progress Report<sup>16</sup>. A new specification, GMS-29100-611 has been issued for the generator to go into the dry agent disseminator because of the modified exterior.

# 7.1.2 Dry Nitrogen System

Dry nitrogen is used in the disseminator to fluidize the disaggregated powder and transport it out of the store through the discharge tube into the slip stream. The nitrogen system schematic drawing is presented in Figure 7.2. The system consists of a high-pressure storage vessel, manual and solenoid shut-off valves, a pressure regulator, a critical flow orifice, a manifold check valve and a manifold with high velocity jets. These basic components are supplemented by pressure gauges, relief valves, pressure switches, etc. required for safe control of the system. Following is a description of the system as it will function in the disseminator.

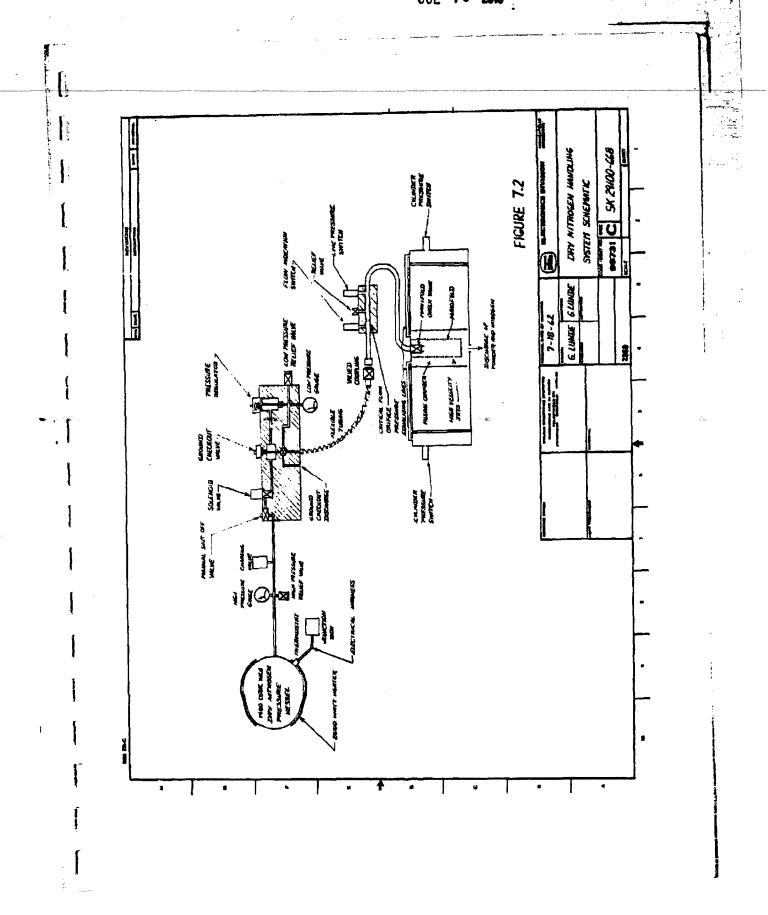
Dry nitrogen is introduced into the pressure vessel through the charging valve and stored at 3000 psi pressure. The manual shut-off valve is closed during charging and storage to insure that no leakage will occur. During pre-flight preparation of the disseminator the manual valve is opened and the solenoid valve is used to stop and start the flow of nitrogen during operation of the disseminator.

The rate of flow of nitrogen to be used during operation is determined by adjusting the pressure regulator to be predetermined pressure before take-off. This controlled pressure in conjunction with the critical flow orifice will give a constant rate of mass flow which is independent of the downstream pressure.

The pressure regulator is adjusted while nitrogen is flowing. A supply of nitrogen is introduced into the checkout valve from an external supply, passes through the pressure regulator, and is exhausted to the atmosphere through the ground checkout discharge. This method of adjustment does not bleed nitrogen out of the system's pressure vessel and will not build up pressure in the inner tank assembly.

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When the nitrogen is released from the pressure vessel, isentropic expansion will cause significant cooling. Under certain conceivable conditions this temperature drop could cause variations of mass flow rate of from 10 to 20 percent. If this problem should arise it can be corrected by adding heat to the gas. For these reasons a heater jacket, laced to the pressure vessel, has been added to the system. If actual test experience shows that the heater is unnecessary it can easily be removed from the system.

A flow indication pressure switch is mounted in the manifold upstream of the critical flow orifice to show whether or not the solenoid valve is open and sufficient pressure is available for proper flow.

The cylinder pressure switches operate when the cylinder pressure exceeds the normal operating level. This would happen if the orifice which discharges the mixture of powder and nitrogen became plugged. The switches will actuate a relay to shut off the gas solenoid valve.

The line pressure switch is used as a safety back-up for the cylinder pressure switches. Its actuation point will be set below the relief valve setting.

The manifold check valve allows free flow of nitrogen into the cylinder but prevents back flow so that the nitrogen system will not become contaminated by agent entering through the manifold.

The pressure equalizing lines allow flow from the mixing chamber to the spaces behind the pistons so that the pistons will not work against a gas pressure differential.

The nitrogen flows through the manifold and out into the mixing chamber via the high velocity jets. The jets will give a velocity of from 150 to 350 ft/sec depending on the flows and the size of the jets used. This velocity will cause swirls in the mixing chamber and will tend to cause good mixing of powder and gas as the powder comes off the disaggregator cutters.



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The nitrogen pressure vessel can store a total of 12.65 pounds or 175 sec of nitrogen at 3000 psi and 70°F. The amount available for mixing with the powder is 9.62 pounds (or 2.75 percent of the 350-pound powder capacity) because of allowances which must be made for changes of flow rate with temperature change, errors in pressure measurement and regulation, and a requirement for a residual pressure to be maintained in the vessel.

With the regulated pressure range of 30 to 100 psig on the fixed orifice, the available range of flow rates is 5.7 scfm minimum to 14.6 scfm maximum. Figure 7.3 shows how this range of nitrogen flow rate combines with the powder feed rate to give various ratios of nitrogen to powder flow. The ratio can be kept at 2.75 percent up to powder feed rates of 38 lb/min by increasing the gas flow as the powder flow is increased. At feed rates above 38 lb/min the maximum flow rate of 14.6 scfm governs. The ratio then decreases until the minimum percent is 1.76 at a feed rate of 60 lb/min.

If the gas-to-powder ratio is allowed to vary from 2.75 down to one percent, only two nitrogen flow rates are needed, 5.7 and 14.6 scfm.

A preliminary study of the standard practices employed in charging high-pressure vessels indicates that there are two practical methods of charging the pressure vessel with dry nitrogen for this application. One is to compress the gas from standard 2200 psi bottles directly into the pressure vessel with a portable compressor. The second method is to order 6000 psi bottles and use these to top off the vessel after filling part way with the readily attainable 2200 psi bottles.

The relative merits of these methods are now being investigated and the necessary facilities will be made available for laboratory and field operation of the nitrogen system.



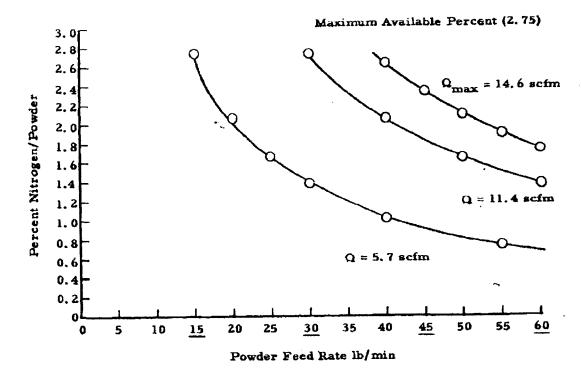


Figure 7.3 Ratio of Nitrogen-to-Powder Flow Rates Obtained from Powder Feed Rate versus Available Nitrogen Flow Rates (Q)

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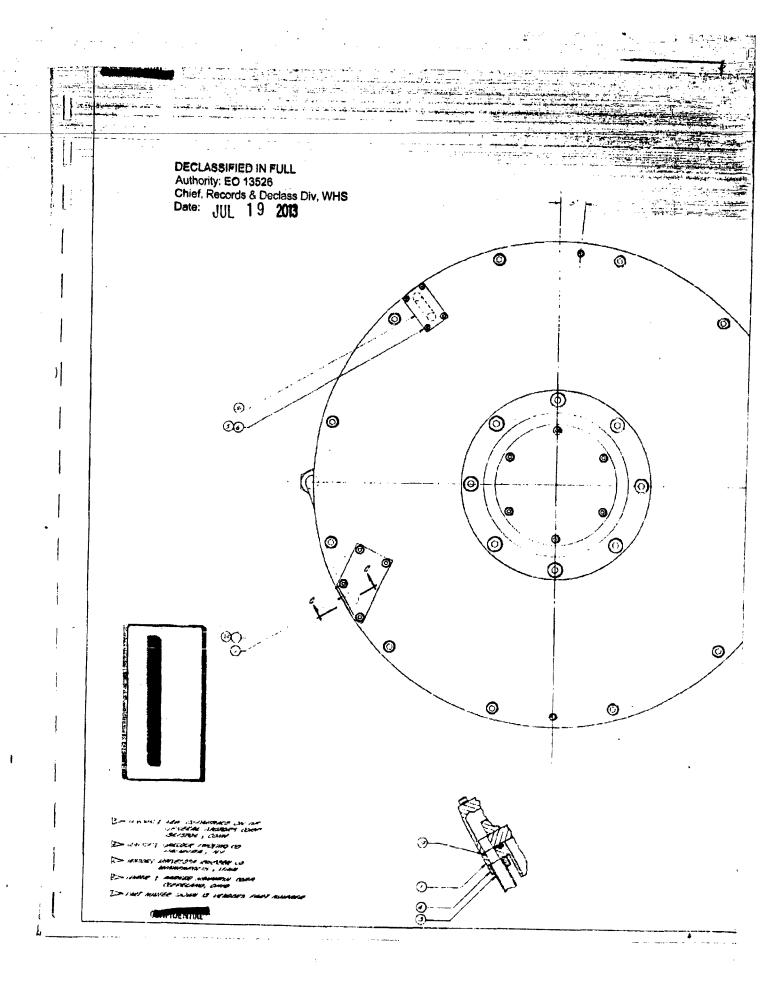
#### 7.1.3 Center Section

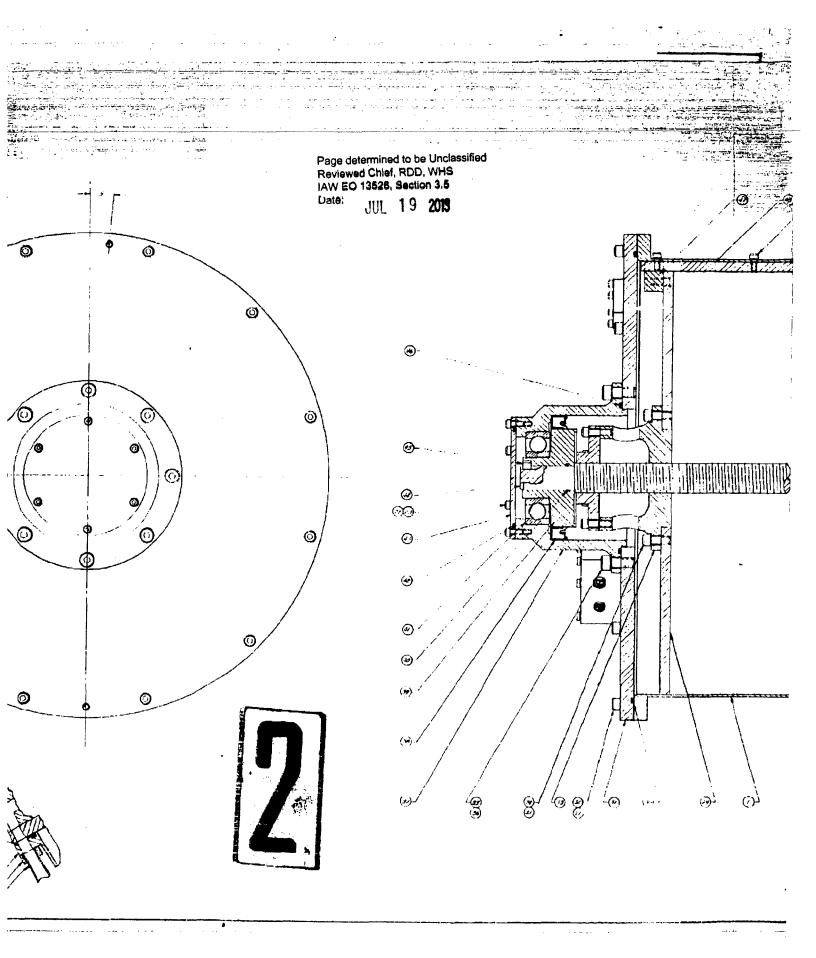
The center section of the store is the basic disseminator. It extends from station 35.5 to station 131.75. It is actually somewhat longer than the length arrived at by subtracting the nose and tail sections from the overall length because the ends of the inner tank assembly extend beyond the stations where the nose and tail joints are made. The construction of this center section will be similar to the liquid agent disseminating store in that the space between the inner cylinder and the external shell will be filled with a foamed-in-place rigid plastic. A strong-back and stiffening rings will be used to satisfy structural requirements. Nitrogen lines and electrical conduits will be placed between the inner and outer shells.

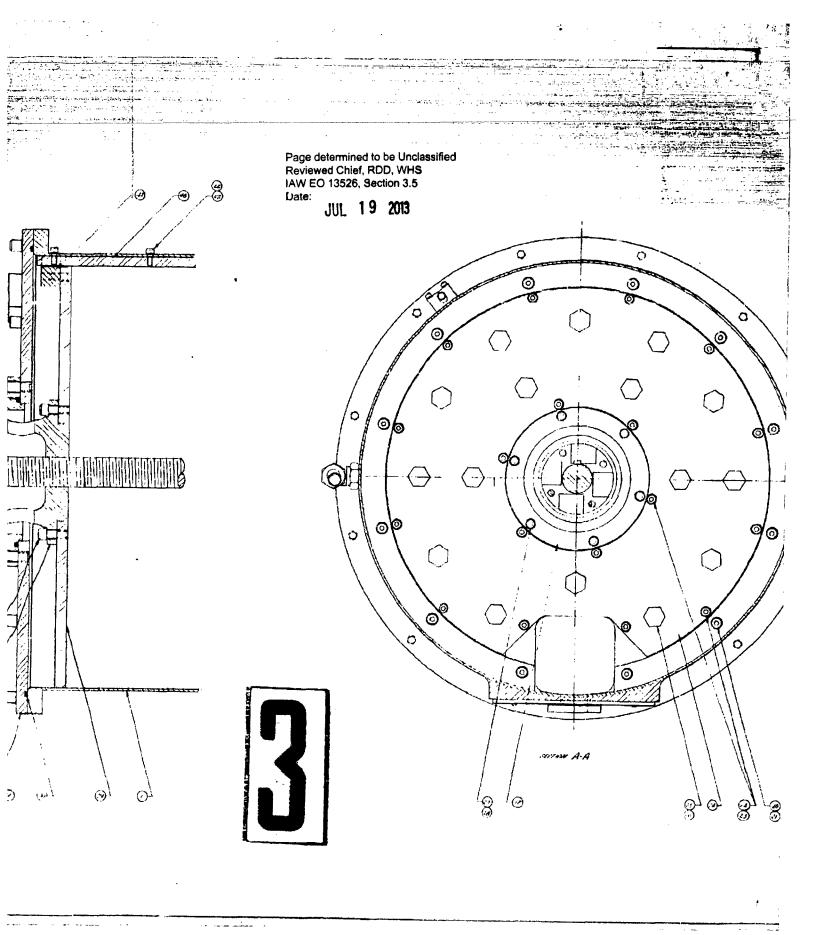
The inner tank assembly consists of a cylinder with removable end plates, a drive screw running the length of the cylinder, two pistons with threaded hubs riding the drive screw, and a disaggregator with cutters in disks keyed to the drive screw at the center. The design of this assembly will be similar to the second experimental model shown in Figure 7.4 and described in Section 7.2. The essential differences in the airborne model as now envisioned will be that a one-piece cylinder with a removable gas manifold will be used to eliminate the center joint shown in the experimental unit, and double O-ring seals will be used on the end plates and all other attachments to the cylinder.

The inner cylinder is 83.25 inches long and 16.5 inches inside diameter. A volume of 9.1 cubic feet is available within the cylinder for containing compacted dry agent material.

The fluidized powder will be discharged through a short tube extending from the bottom of the center section. An NACA study <sup>17</sup> of discharge tubes of this type showed that in order to obtain good separation of the discharge flow from the boundary layer of the store, the tube should be housed by an air foil capped with an air-flow control plate. In the case of the disseminator it is necessary that the tube shroud be large enough to house the valve mechanism and, consequently, a tapered eliptical shroud is considered to be the best compromise design. Such a shroud is shown in Figure 7.1.

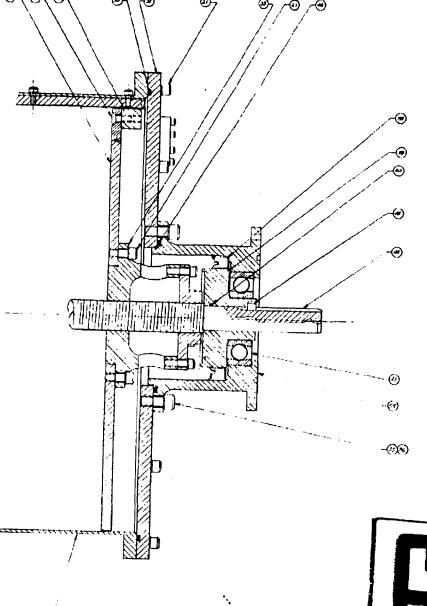






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# 7.1.4 Rotary Actuator Assembly

The requirements of the rotary actuator for driving the feeding mechanism, as outlined in the Seventh Quarterly Progress Report<sup>18</sup> and repeated below, were submitted to a number of potential vendors but none could furnish a satisfactory item without resorting to a design and development program. Since General Mills, Inc. has had considerable experience in developing equipment of this type, a decision was made to have our personnel proceed with the design and fabrication of the actuator.

The requirements of the actuator are:

- Output Speeds. 12, 18, 24, 36 and 48 rpm in either direction. (These speeds are changed from those given in the referenced report.)
- Output Torque. 2500 pound-inches in either direction at the above speeds.
- 3) Maximum Allowable Overhung Shaft Load. 1500 pounds.
- Maximum Allowable Inward Thrust Load on Shaft. 2000 pounds.
- Maximum Allowable Outward Thrust Load on Shaft. 2000 pounds.
- 6) Duty Cycle. Continuous for periods up to 1/2 hour.
- 7) Life. 200 hours.
- 8) Operating Temperature. 160°F to -65°F.
- 9) Operating Altitude. Sea level to 15,000 feet.
- 10) Acceleration. 10 g's in any direction.
- Vibration. 5 to 500 cps at 0.036-inch double amplitude or +10 g whichever is the lower value.
- 12) Input Electrical Characteristics. 400 cycle, 200 volt, 3 phase, ac.
- 13) Connectors. Water-tight connector at cable entrance to actuator housing.

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During this reporting period the design was essentially completed and all major items were released for fabrication or procurement. As the design progressed, it was necessary to modify some of the characteristics so that they now differ from the description presented in the Seventh Quarterly Progress Report 18.

The electric motor which has been ordered from the Westinghouse Electric Corporation is a 400 cps, 3-phase, 200-volt, 5600-rpm motor with a rated torque capacity of 22 pound-inches. The speed-selector portion of the actuator will still provide for five driving speeds as given in Item (1) above. At all but the highest speed the speed-changing gearing will all provide some speed reduction. The highest ratio is actually a step-up to about 117 percent of input speed.

The major speed reduction still occurs in the fixed-ratio speed-reducing part of the actuator which is a series three-stage planetary gear unit. The reduction ratio of the first and second stages is 4.75 to 1 each, and that of the third stage is 6 to 1, giving an over-all reduction ratio of 135.375 to 1. The input in each stage is at the sun gear and the output at the planet carrier or its extension. The incoming torque is received through a safety clutch which is designed to slip when the force exceeds a given amount.

The approximate size of the actuator package is now 12 inches maximum diameter and 21 inches length. Slightly more than half of the unit has a diameter of about 5 inches. The weight has been calculated at 65 pounds. The actuator assembly will have two mounting flanges for securing the unit in the tail section of the stors. The principal mounting of the actuator is at the forward bulkhead of the motor housing and the secondary mounting is the flange portion of the housing at the forward or output end of the planetary speed-reducing unit.

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# 7.2 Fabrication of the Second Experimental Unit

The design for the airborns dry agent disseminating store which is evolving from design studies and laboratory experiments will differ in some respects from the full-scale feeder which has been used thus far in this phase of the program. The same basic concepts will be employed but, in general, the experimental feeder design is not directly applicable to the airborne store. Therefore, a second experimental model is being fabricated to test the design to be employed in the airborne store. This new experimental unit is shown in Figure 7.4 (GMI Dwg SK-29100-778). The more important changes in this model as compared with the first experimental feeder are discussed below.

- 1) The cylinder dimensions are reduced to 83.25 inches long by 16.5 inches inside diameter. These dimensions are those of the inner tank to be incorporated in the airborne version. Stainless steel, type 304 ATSI, is utilized rather than aluminum as in the first generation model.
- 2) The drive screw is machined as one piece rather than two as on the first generation model. This eliminates the problem of joining the two screws together when assembling the unit. The disaggregator is keyed to the drive screw to permit easy removal of the screw for cleaning and maintenance.
- 3) Ball bearings for radial and thrust support are again used at the ends of the screw but additional bearings are mounted at the center of the cylinder to support the disaggregator and reduce friction at this point. The center support structure has been incorporated in the aerating ring structure to obtain a compact arrangement.
- 4) The orifice plates forming the sides of the aerating ring or manifold have removable nozzles for directing the gas which mixes with the powder and causes it to flow out of the unit.
- The piston hubs are elongated to provide better support on the drive screw.
- 5) The experimental unit is designed for loading from the ends, using a special loading tube, as is planned for the airborne version.

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Controls for the gas system are not shown in Figure 7.4 but the following items will be installed on the experimental unit to test for performance:

- Pressure switches to indicate abnormally high pressure within the cylinder at the center and at the ends behind the pistons.
- A check valve at the entrance to the aerating ring to prevent back flow.
- 3) A pressure switch to indicate abnormally high pressure in the gas line entering the aerating ring.
- 4) A pressure relief valve in this line.
- A fixed critical flow orifice ahead of the pressure relief valve.
- 6) A pressure regulator in the gas supply line.

This experimental unit will be tested with the same facilities which have been used to obtain data with the first experimental unit.

# 7.3 Fabrication of Loading Equipment for Use With the Second Experimental Unit

Loading of the disseminator with compacted dry agent will require auxiliary equipment to compress the finely-divided bulk solids into cylindrical packages of the required length, diameter and density. Special equipment may also be necessary to transfer the compacted powder from the press to the disseminator. In order to study this aspect of the program, a hydraulic press and a loading fixture have been designed for use with the second experimental unit. Fabrication of this equipment was started during this reporting period.

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### 7. 3. 1 Hydraulic Press

The hand press employed in loading the full-scale feeder with compacted powder is not suitable for producing densities greater than approximately 0.45 grams per cc. Since the objective is to investigate disseminator performance with compaction densities up to 0.6 grams per cc, it will be necessary to have a press capable of producing the total force associated with this density. Gonsequently, a simple hydraulic press is being fabricated which will be capable of exerting a force of 8300 pounds. The compaction ram will be attached to a hydraulic cylinder having a 38-inch stroke. Hydraulic controls will be provided to enable the operator to vary the rate of piston travel and to adjust the piston force.

#### 7.3.2 Loading Fixture

A loading fixture is being fabricated which consists of a loading tube and a manual lift truck. The loading tube will be positioned in the hydraulic press for filling and compacting of the powder. The lift truck will be used to raise the filled loading tube and rotating it to a horizontal position in line with the disseminator. The loading tube will then be attached to the end of the inner tank of the disseminator and the compacted material will be pushed into the disseminator using air pressure to operate the loading tube. The loading fixture will then be removed and the piston and end plate of the disseminator will be installed.

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# 8. TESTING OF THE LIQUID AGENT DISSEMINATING STORE

The liquid agent disseminating store, which was described in the Seventh Quarterly Progress Report<sup>19</sup>, has been subjected to a series of structural and functional tests in the laboratory and in the field. Laboratory structural tests were conducted at Fletcher Aviation Company, El Monte, California using two test units which were fabricated for that purpose. The third unit, which is a complete developmental model, was tested in the laboratory at General Mills, Inc. and, subsequently, flight tested on an A4D-1 airplane at the Naval Air Test Center, Patuxent River, Maryland. The disseminator has met the requirements of the various tests with a very high degree of success. An assembly drawing showing the complete unit is included as Appendix A of this report.

A report (see Appendix B) of the structural test program is included with this progress report and is discussed briefly below. The results of the test work conducted at General Mills, Inc. will be presented in a final engineering report which is being prepared. The Naval Air Test Center is submitting an official report on the flight tests. A short description of the flight test project is presented in paragraph 8.2 following.

# 8. 1 Structural Testing at Fletcher Aviation Company. El Monte, California

Appendix B is Fletcher Aviation Company Report No. 43.286, "Qualification Tests, General Mills Tank Assembly" covering the structural testing conducted by Fletcher on two units fabricated for this purpose. The test models were structurally similar to the delivered unit with the exception that components such as the turbine, pump, actuator, etc., were simulated by means of dummy units having the same weight, center of gravity, and attachment provisions.

The tests were conducted in general accordance with Specification MIL-7378A, "Tanks, Fuel, Aircraft, External, Auxiliary, Removable." Following is a list of the test performed:

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- Examination of product for conformance with drawings and for quality of workmanship.
- 2) Determination of weight and center of gravity locations.
- 3) Determination of tank capacity.
- 4) Slosh-and-vibration test.
- 5) Leakage test.
- 6) Ground ejection test.
- 7) Static structural test.

The disseminator successfully met the requirements of the various tests. In order to pass the slosh-and-vibration test it was necessary to stop the test after it had been in progress for 17-1/2 hours and repair a crack in the skin and add reinforcement strips as described in pages 4.7 through 4.12 of Appendix B. After this modification the tank successfully withstood an additional 25 hours of slosh-and-vibration testing.

During the repair, the inner tank was inspected. It was discovered that the buna rubber lining of the inner Fiberglas tank had separated from the tank and was torn in several places (see page 4.11 of Appendix A). In addition, the two sati-slosh baffles were intact but had broken free and were lying on the bottom of the tank. The lining and the bulkheads were removed before the test was repeated.

When this Fiberglas tank was delivered to Fletcher for incorporation into the assembly, it was known that the bond between the liner and the tank proper was inferior. The liner was already separated from the tank in places. Since this liner was used to facilitate release of the tank from the mold during fabrication and is not required for structural or leakage purposes, this fault was not considered to be important.

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# 8. 2 Flight Tests of Liquid Agent Disseminating Store at NATC, Patuxent River, Maryland

Through arrangements with the Bureau of Naval Weapons it became possible to conduct flight tests with the liquid agent disseminator using an A4D airplane. The unit was shipped to the Naval Air Test Center, Patuxent River, Maryland where a series of flight tests were conducted by the Weapon Systems Test Division on 16, 17, 18 May, 1962. The WST Division was directed to prepare a final report covering these tests. The following remarks concerning the flight tests are based on observations made by General Mills, Inc. personnel who participated in the tests and on preliminary verbal reports made by Lt. H. Turk, the test pilot.

The following "detailed Requirements" were listed in Weptask No. RMMO-33-015/201-1/F008-10-005 issued by the Bureau of Naval Weapons for the flight test project. The unit successfully passed all phases of this testing program.

- 1) Perform fit tests with the spray tank suspended from the Aero 7A Bomb Rack of the A4D-1 aircraft.
- Provision for appropriate electrical connectors in the aircraft pylon, if required.
- Perform static functional spray tests using water.
- 4) Perform flight tests to the maximum safe speeds not to exceed the limits of normal flying as set forth in BUWEPS Instruction 3710.0 of 19 October 1960.
- 5) Pilot to observe and report unfavorable conditions during taxing, take-offs, landings, and maneuverability tests (high and low altitude, high and low speed).
- 6) Perform high altitude, low temperature soak followed by functional test with dyed water. Upon landing make a visual check of aircraft and report areas covered with dye. Report temperature and duration of cold soak.
- 7) Perform low altitude (300 to 500 feet) high-speed functional test with dyed water. Upon landing make a visual check of the aircraft and report areas covered with dye.



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- 8) Provide for camera coverage of functional test items (3),(6) and (7) above.
- Furnish twelve photographic copies of the GMI spray tank and aircraft installation to BUWEPS (Code RMMO-334).

The photographs in Figures 8.1 and 8.2 show the disseminator mounted on the Aero 7A bomb rack on the fuselage centerline station of the A4D-1 airplane. Electrical connections to the store were easily made by installing a cable running from the pylon to a junction box in the fuselage just forward of the pylon. The cockpit control panel was installed in a position used for such auxiliary equipment and required no airplane modifications.

The static functional spray tests were conducted with the unit mounted on a bomb rack hanging from a steel frame provided by the Navy. A ground power source was used to operate the disseminator. Figure 8.3 shows water spraying from the booms during this test.

The flight tests were conducted in three flights. The pilot reported no unfavorable conditions due to the disseminator during these flights and found the control panel to be entirely satisfactory. The unit functioned properly at all times.

The high-altitude, low-temperature test was conducted on the second flight in which two auxiliary 300-gallon fuel tanks were installed at the wing stations to obtain the desired flight duration. Two minimum-maximum thermometers mounted in the aft section of the disseminator for this test indicated a temperature range from 73°F to 80°F.

Maneuvers resulting in 5 "g" loading were performed on the third flight. Following these maneuvers, several low-altitude dissemination runs were made within sight of an observation tower so that motion pictures could be made with a telephoto lens. The dissemination process was visible to the naked eye and the aerosol was observed to trail out from the booms of the disseminator in a clearly defined band that did not diffuse until it was well aft of the airplane.

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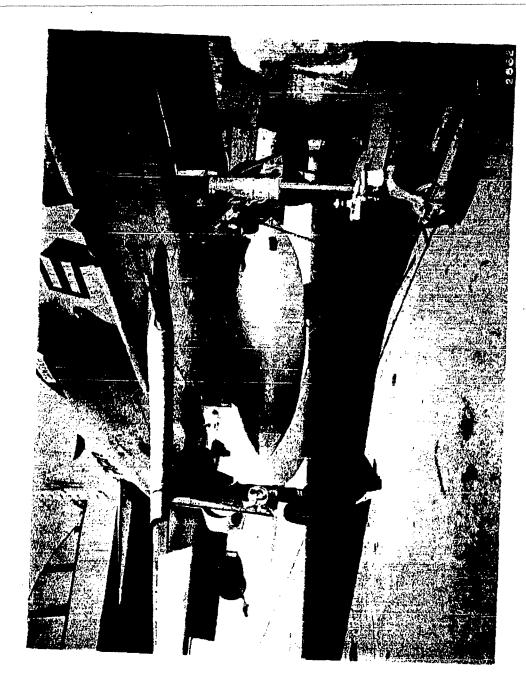
Figure 8.1 GMI Liquid Agent Disseminating Store Mounted on Aero 7A Bomb Rack on A4D-1 Airplane - Side View

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GMI Liquid Agent Disseminating Store Mounted on Aero 7A Bomb Rack on A4D-1 Airplane - Front View Figure 8.2

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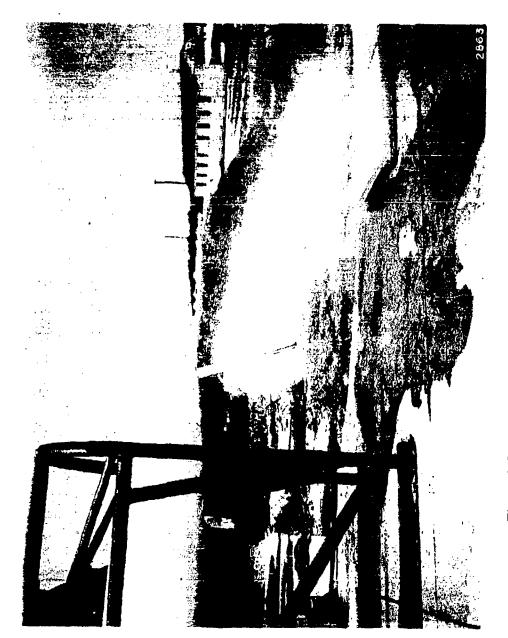


Figure 8.3 GMI Liquid Agent Disseminating Store Spraying Water During Static Functional Test

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For each of the three flights the disseminator was filled with dyed water to aid in photography and to provide a tracer for studying contamination of the aircraft. Both methylene blue and uranine (sodium fluorescein) were used at a concentration of approximately 0.25 percent each. The methylene blue was planned to produce evidence of contamination visible under normal light and the uranine under ultraviolet. A very careful examination of the airplane was made after each flight and no evidence of contamination was found. In fact, the only areas on the store itself which were contaminated were the booms, the boom wells, and the exterior surfaces immediately adjacent to the boom wells.

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#### 9. SUMMARY AND CONCLUSIONS

During this reporting period, significant work was accomplished in our research and development program on the dissemination of solid and liquid BW agents. Progress in each of seven areas of effort is summarized below with the pertinent section of this report indicated at the end of the paragraph.

Data obtained with an improved piston-cylinder compaction apparatus have resulted in an empirical formula of the form  $\sigma = K(\frac{1}{A})^{-1}$  relating stress, of, and density, p, of a compacted powder. Values for K and r were determined for tale, saccharın, powdered sugar, powdered milk and cornstarch. Tests with the hydrostatic compaction apparatus yielded results in agreement with the improved piston-cylinder compaction apparatus, but the hydrostatic apparatus proved to be difficult to use. Measurements at high stress levels with the above powders indicate that considerably less elastic energy is stored in a compacted powder bed than was first assumed. It has been found that very little elastic recovery occurs as stress is reduced at high stress levels. The triaxial shear tests have thus far proven successful only with relatively highly compacted samples and low (2 psi or less) radial pressures. Additional data were obtained using the bulk tensile strength apparatus with zinc cadmium sulfide which indicates that total sample length and length of time of application of compressive load when preparing the sample are important considerations. Bulk density investigations have produced data showing a decrease in density with increased radial distance from the axis of a sample compacted in a cylinder. Using the sliding disk method to measure shear strength as a function of compressive stress it has been observed that the relationship remains unchanged as humidity is increased from 2 to 32 percent but marked changes were observed at 46 and 69 percent (Section 2).

The aerophilometer has been operated while studying experimental techniques and determining operating parameters essential to carrying out the program on stability properties of aerosols. A mathematical analysis has produced theoretical expressions which, as shown by experiments,

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have a considerable degree of validity and have proven very useful in interpreting the light-scattering data. The decrease in scattered light with time has been recorded for zerosols of tale, saccharin, powdered milk and K-ferric oxide using the fan to produce turbulence. Erratic behavior was observed when tranquil conditions were employed. Apparently, "clouds" form in the zerosol and drift randomly through the light-scattering area (Section 3).

An experiment has shown that <u>Sm</u> powder being used in trials designed to measure the degree of heat inactivation were invariably contaminated with <u>Bg</u> colonies. A series of trials have demonstrated that neither coating with <u>Cab-O-Sil</u> nor compaction to 0.62 grams per cc density has any significant or deleterious immediate effects on viability of <u>Sm</u> (Section 4).

A program was conducted at Fort Detrick under Technical Evaluation Division Test No. 62-TE-1602; MD Division No. 1927, using the 40-foot diameter test sphere to evaluate the General Mills GMI-3 fixture and wind tunnel when used for generating aerosols of dry Sm and dry P. tularensis. When the Technical Evaluation Division furnishes the statistical analysis of data, it will be possible to report on the effects of compaction and subsequent aerodynamic breakup during dissemination on the viability of dry agents. Preliminary examination of the data indicates that good recovery factors were obtained (Section 5).

The full-scale experimental feeder for use with compacted dry agent simulants has been operated successfully over material flow rates ranging from 20 to 49 lb/min. It has been demonstrated that reasonably low gas flows are sufficient to fluidize the powder and carry it out through a discharge tube. Although the feeder has been operated satisfactorily with gas flow as low as 3 scfm, tests have shown that best performance has resulted when the rate was approximately 6 scfm. Torque and power required to drive the feeder have both been well below the limits established in the design studies for an airborne disseminator based on the basic principles employed in the full-scale feeder (Section 6).

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The design studies on a dry BW agent disseminating store have progressed to the stage where major components are now well defined and a preliminary general arrangement has been prepared. The store will have the same external shell as a standard 150-gallon auxiliary fuel tank and will have provisions for both 14- and 30-inch spacing of the mounting lugs. The agent will be contained within a tank assembly forming an integral part of the center section of the store. The air turbine generator and the gas supply vessel will be housed in the nose section, and the rotary actuator in the tail. Both nose and tail sections will be attached to the center section with bayonet-type joints. The dry agent will be discharged through a tube projecting below the store sufficiently far so that the material is injected into the slipstream beyond the boundary layer. An experimental version of the inner tank assembly is being fabricated for use in laboratory tests of the engineering design (Section 7).

The liquid agent disseminating store was flight tested on an A4D-1 airplane by the Weapons System Test Division at the Naval Air Test Center,
Patuxent River, Maryland. Dyed water was successfully disseminated
under various conditions and no trace of dye could be found on the airplane
after it returned to the ground. The pilot reported no unfavorable conditions
of flight and there was no damage to the store during maneuvers resulting
in 5 "g" loading and speeds up to the maximum safe limits of normal flying
for the A4D-1 airplane (Section 8).

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- 15) ---- Report No. 2300, op. cit., pp. 8-1 to 8-5.
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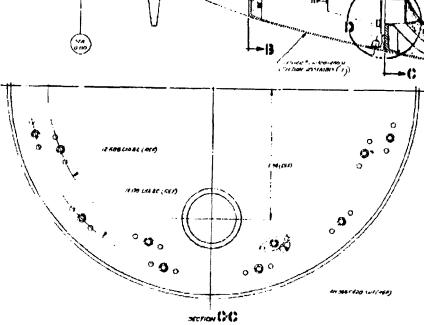
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## APPENDIX A

ASSEMBLY DRAWING
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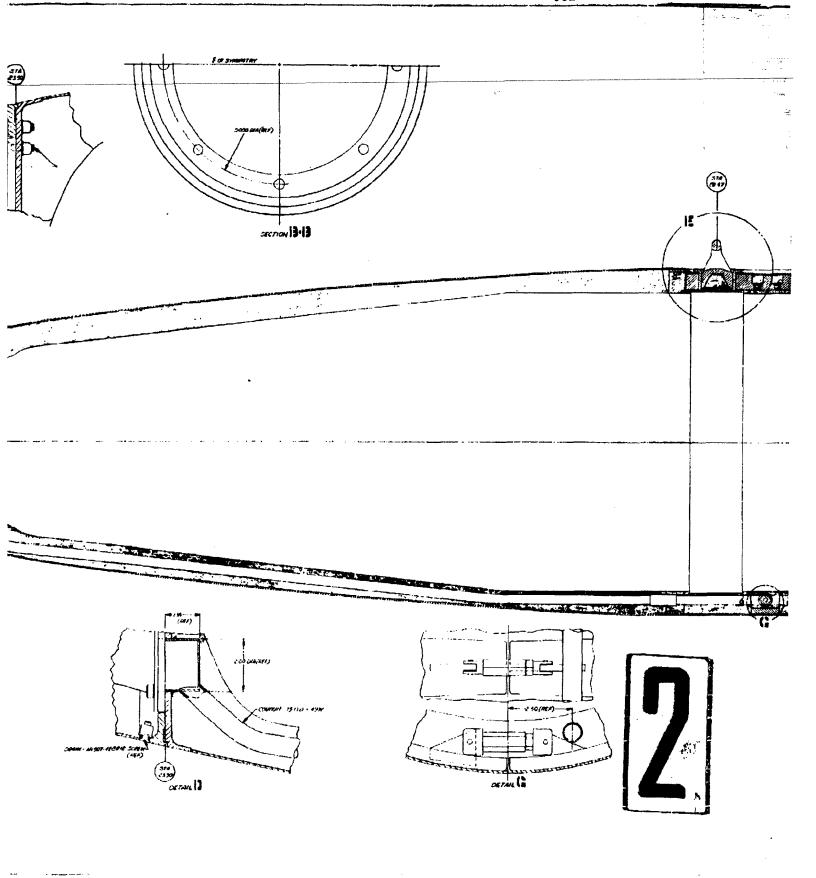
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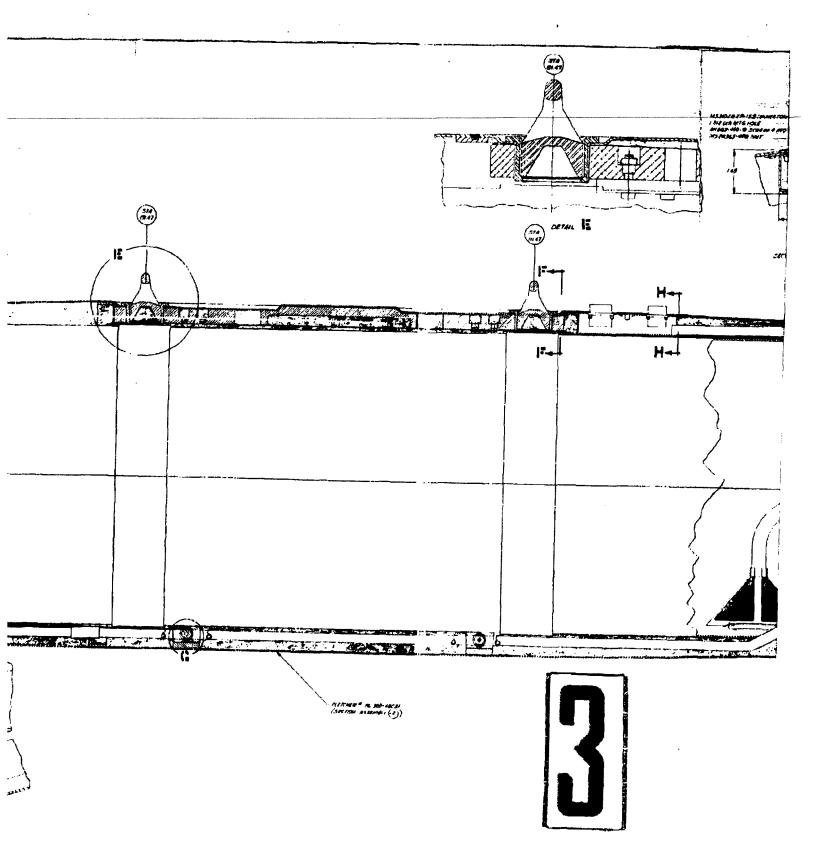


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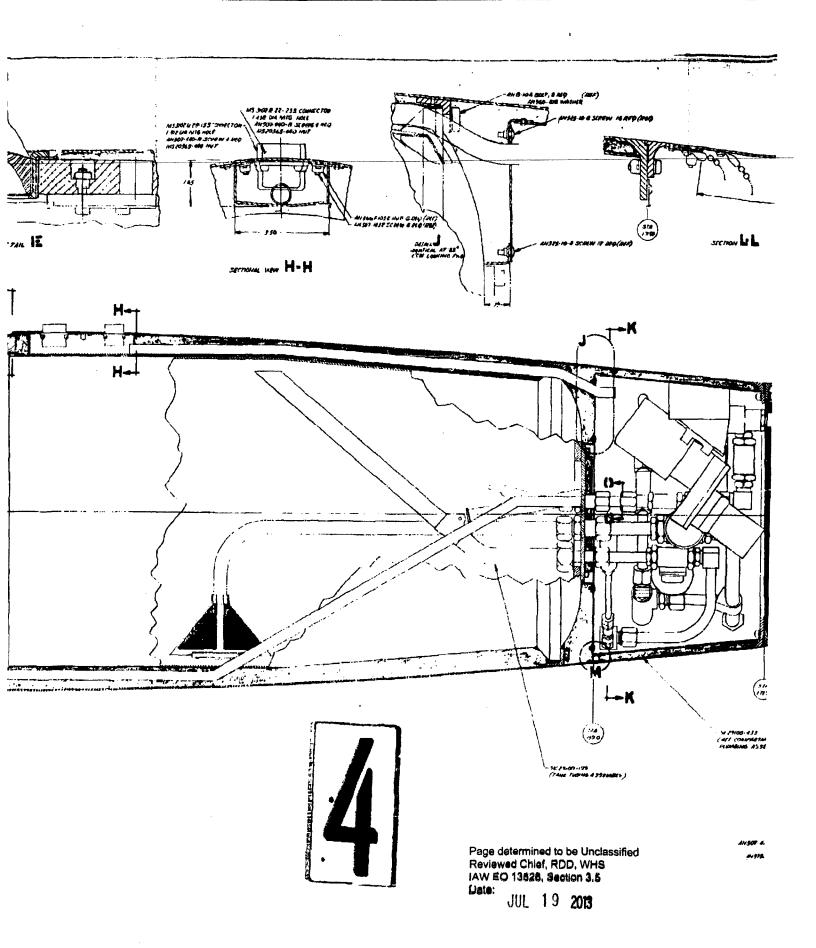
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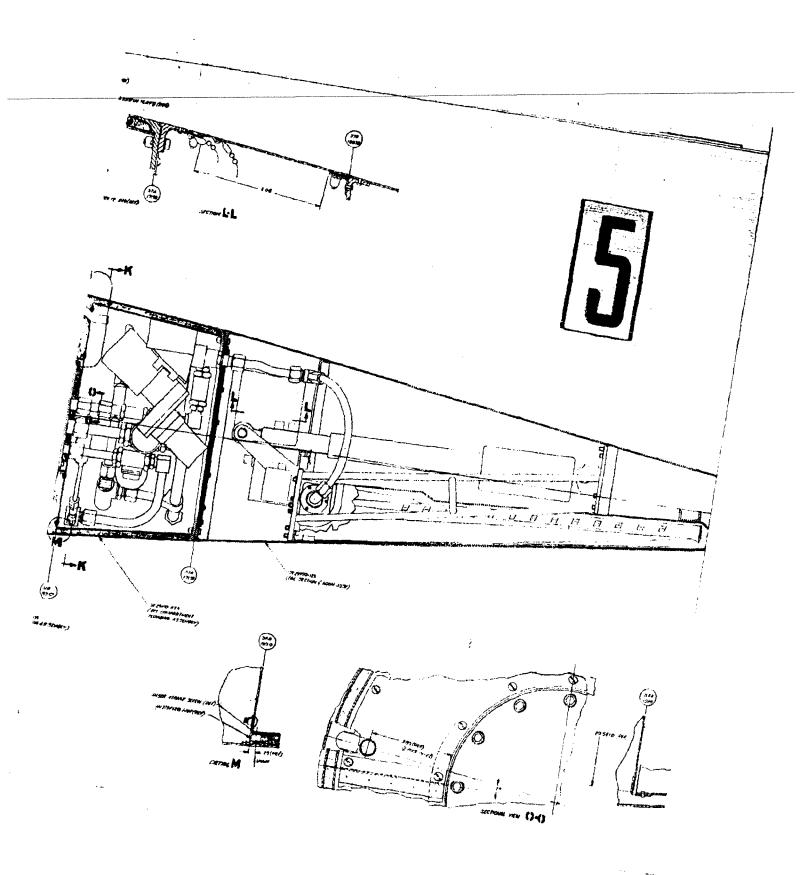
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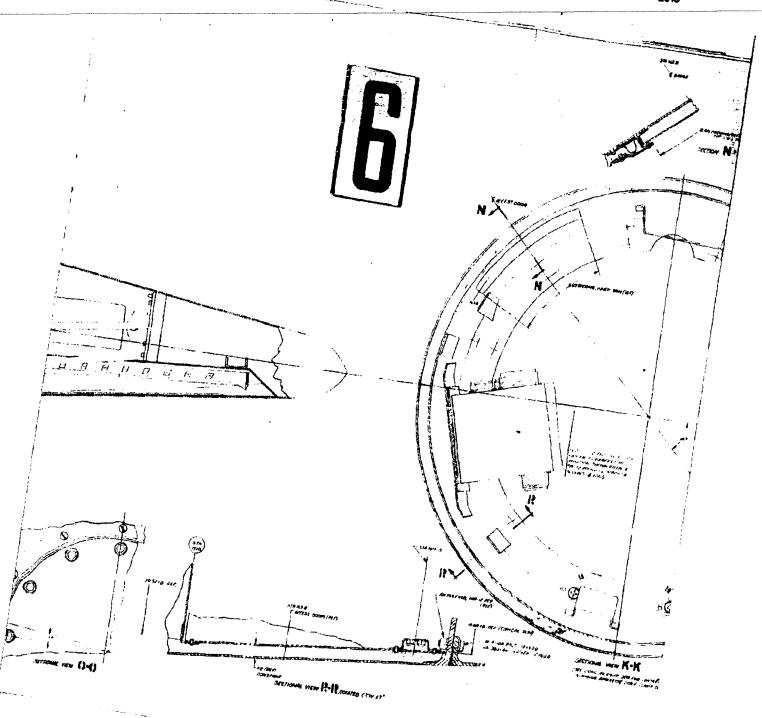




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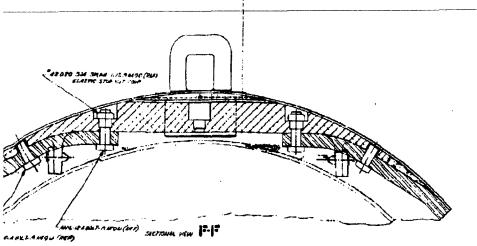


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#### APPENDIX B

FLETCHER AVIATION COMPANY REPORT NO. 43.286

"QUALIFICATION TESTS, GENERAL MILLS TANK ASSEMBLY" REPORT NO. 43.286

DATED

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# FLETCHER aviation corporation

QUALIFICATION TESTS GENERAL MILLS TARK ASSY

APPROVED

MODEL

26-300

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#### INTRODUCTION

This report covers a portion of the qualification tests conducted on General Mills (G.M.I.) special tank assembly, Part No. SK29100-026. The test assemblies are identical to the final assembly except that operational components such as the ram air turbine, generator, pump, valves actuator, etc., were models having the same weights and center of gravity locations. Further qualification testing will be conducted by G.M.I. to substantiate the operational capabilities of the complete assembly.

This document presents the qualification testing conducted in accordance with the requirements of specification MIL-T-7378A.

Qualification tests described herein were conducted at the Fletcher Aviation Company test facility.

#### REFERENCES:

Specification MIL-T-7378A - Tanks, Fuel Aircraft,
External Auxiliary,
Removable

G.M.I. Specification - SK29100-026

G.M.I. Specification - External Removable Tank
GMS-29100-026 - Assy for G.M.I. Electronics
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F.A.C. Drawing 26-300-48031 - Tank Assembly, G.M.I. Specification

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#### ADMINISTRATIVE DATA

#### PURPOSE OF TEST:

To determine that the assembly conforms to the applicable drawings.

## MANUFACTURER:

Fletcher Aviation Company

MANUFACTURER'S MODEL NO:

26-300

ASSEMBLY DRAWING:

26-300-48031

QUANTITY OF ITEMS,

One

SECURITY CLASSIFICATION:

None

TEST DATE:

1-30-62

TEST CONDUCTED BY:

Wayne Callahan

DISPOSITION OF SPECIMEL:

Use for Weight Test

#### ABSTRACT:

The assembly successfully met all of the requirements of the test. Rubber liner of inner tank was not tight in several areas.

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#### FACTUAL DATA

#### REQUIREMENTS:

- 1. That the assembly and accessible components thereof conform to their applicable engineering drawings.
- 2. That the assembly and all components conform to aircraft quality standards for workmanship.

#### THET BOULPMENT:

 Standard inspection tools; linear scales, micrometers, calipers.

#### TEST PROCEDURE:

- Inspect the complete assembly for general conformance to F.A.C. drawing no. 26-300-48031.
- Remove access doors and inspect all compartments for metal chips, filings, or other foreign material.
- Inspect all mating components and access doors for fit, sealing capabilities and general workmanship.
- Inspect tank and inner compartment surfaces for evidence of damage or undue abrasion.
- Inspect for loose bolts, rivets, or other fastening devices.
- Inspect for parts not treated for corrosion registance.
- 7. Inspect for misalignment of mating components.
- 8. Remove inner tank cover plate and inspect inner tank for general cleanliness, worksanship, and conformance to G.M.I. requirements.
- 9. Generally inspect simulated components (G.M.I. furnished) for fit and security of installation.
- 10. Install doors and inspect entire tank contour for surface irregularities.

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#### ADMINISTRATIVE DATA

PURPOSE OF TEST:

The second second

To determine the weight and location of the center of gravity.

MANUFACTURER:

Flatcher Aviation company

MANUFACTURER'S MODEL NO:

26-300

ASSEMBLY DRAWING:

26-300-48031

QUANTITY OF ITEMS:

One (1)

SECURITY CLASSIFICATION:

None

TEST DATE:

2-1-62

TEST CONDUCTED BY:

Wayne Callahan

DISPOSITION OF SPECIFUL:

Use for Capacity Test

ABSTRACT:

The assembly successfully met all of the requirements of the test.

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#### FACTUAL DATA

#### REQUIREMENT:

- 1. To determine the total weight of the empty assembly.
- To determine the center of gravity locations to the requirements of the detail specifications for the tank.

#### TRET SOUI PHENT:

- 1. Two platform scales.
- 2. Tank cradle.
- 3. Rocker bar.
- 4. Water meter.

#### TEST PROCEDURE:

- 1. The cradle is weighed.
- The tank and cradle are weighed and the cradle weight is deducted from the total weight.
- 3. The cradle is balanced on the rocker bar.
- The tank is placed on the cradle in the "balance" position. The c.g. station is recorded.
- 5. Water is metered into the tank to 1/4 of the rated capacity.
- 6. The force and moment arm required to balance the tank at a range of tank attitudes from 5° nose down up to and including 30° nose up.
- Items 5 and 6 are repeated for the 1/2, 3/4 and full capacity.
- 8. The c.g. for each condition is determined from the total weight and the balancing force.

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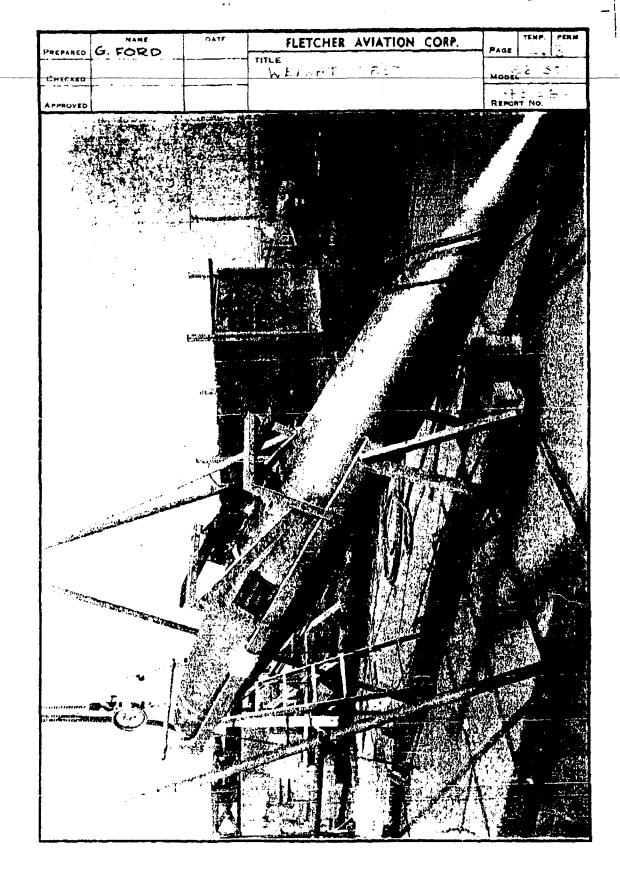
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			ADMINISTRATIVE DATA	<b>(3)</b>	
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	PARPARED W.Callahan	1-5-62	FLETCHER AVIATION COMPANY	PAGE	TEMP.	3.1
1	CHFCKED	ļ	Capacity test General Mills Tank Assy	MODEL		
-	APPROVED.	· !	GENERAL RILLS TARK ABSY	REPOR	43.2	≀86

#### PACTUAL DATA

#### TEST REQUIREMENTS:

To determine total volume of the foamed in place inner tank assembly.

#### TEST EQUIPMENT:

- 1. Bowser Exactometer.
- 2. Suitable graduated container.

## TEST PROCEDURE:

- The tank shall be supported at 90° to normal ground attitude with the inner tank opening in the uppermost position.
- 2. The tank shall be filled to the top and the volume recorded.

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W.Callahan	1-5-62	FLETCHER AVIATION COMPANY	PAGE	TEMA
· ;		CAPACITY TEST		
		GENERAL MILLS TANK ASSY	Mode	<u>26-</u> 43.
ED			REPOR	
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TEST RESULTS:	\$			
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	PREPARED	W.Callahan	1-5-62	FLETCHER AVIATION COMPANY	PAGE	78MP.	PRIN.
~	CHECKLO			SLOSH/VIBERTION TEST	Mace	26-3	100
	APPROVED			GENERAL MILLS TANK ASSY	REPOR	43.2 T NO.	86

## ADMINISTRATIVE DATA

(3)

## PURPOSE OF TEST

To demonstrate the tank will withstand vibration and pitching forces encountered in service.

#### MANUPACTURER:

Fletcher Aviation Company

MANUFACTURER'S MODEL NO. :

26-300

ASSEMBLY DRAWING:

26-300-48031

QUALITATY OF ITEMS:

One (1)

SECURITY CLASSIFICATION:

None

TEST DATE:

2-5-62

TEST CONDUCTED BY:

Wayne Callahan

DISPOSITION OF SPECIMEN

Use for Leakage Test

## ABSTRACT:

The assembly successfully met all of the requirements of the test.

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PREPARED W. Callahan 1-6-6	TITE	PAGE	raise.	4.1
CHECKED .	GENERAL MILLS TANK ASSY	Model	43.	-300 2 <b>86</b>

#### FACTUAL DATA

#### REQUIREMENTS:

1. The complete test specimen is mounted in the support fixture on the slosh and vibration machine. The centerline of the tank, so mounted, is a minimum of twenty inches above the slosh axis. The tank booms are in the retracted position.

The test specimen is filled two thirds full with water at ambient temperature and is simultaneously slosh and vibration tested in accordance with the following conditions.

- A. The vibration displacement is a minimum double amplitude of 0.032 average between the top and bottom of the tank and at the supporting rings. The average peak value, at the point of measurement during a thirty second interval, is the value recorded.
- B. The vibration frequency is  $2000^{+0}_{-60}$  C.P.M.
- C. The tank is mounted with the major horizontal axis at 90° to the axis of the shaft of the rocker platform.
- D. The slosh angle is 30° total, approximately 15° on either side of the horizontal position.
- E. The tank is pressurized to 15 P.S.I.G.
- F. The tank is slosh-vibrated for 25 hours at 16 to 20 slosh C.P.M.
- Pollowing the slosh and vibration test, the tank booms are extended with the tank two-thirds full with water at ambient temperature and is :imultaneously slosh and vibration tested in accordance with the following conditions:
  - A. Repeat steps "A" through "E" of the previous test.

PREPARED	W.Callahan	1-6-62	FLETCHER AVIATION COMPANY	Page	TEMP.	4.
CHECKED			SLOSH/VIBRATION TEST GENERAL MILLS TANK ASSY	Model		-300 286
APPROVED		!		REPOR		

## FACTUAL DATA

## ABOUIREMENTS (contd)

- B. The tank is slosh-vibrated for 15 minutes at 16 to 20 slosh CPM.
- 3. The tank is filled with water and vibrated for 10 minutes at vibration displacement specified in "A" of the previous test.

## TEST EQUIPMENT:

- 1. Slosh and vibration machine.
- 2. Tank support fixture.
- 3. Water meter.
- 4. Strobotac or equivalent.
- 5. Vibration meter.
- 6. Vibration pickup.
- 7. Pressure gage.

## TEST PROCEDURE:

- 1. The test assembly is mounted on the slosh and vibration machine by means of a support fixture. The centerline of the assembly is a minimum of 20 inches above the slosh axis.
- 2. The tank is filled with water in the amount corresponding to 2/3 of the rated capacity of the tank in gallons.
- 3. The tank is pressurized to 15 P.S.I.G.

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# (1)

# FACTUAL DATA

## TEST PROCEDURE: (contd)

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- 4. The slosh and vibration machine is put into cyclic pitching of the longitudinal axis from 15° nose up to 15° nose down.
- 5. The motor drive for the accentric weights, producing vibration is activated and brought up to 2000 +0 cpm.
- Vibration readings are taken and the occentric weights adjusted to produce the required values of vibration displacement.
- 7. The slosh rate is checked to be 17 per min.
- 8. The rotation speed of the eccentric weights is measured to be 2000  $^{+0}_{-60}$  R.P.M.
- The air pressure is checked and adjusted if necessary, to 15 P.S.I.G.
- 10. Simultaneous slosh and vibration is continued for 25 hours.
- 11. The tank booms are extended and the tank is simultaneous slosh vibrated for 15 minutes in the configuration described in steps 1 through 9.
- 12. Note the resonant vibration behavior of the booms.
- 13. The tank is then depressurized, filled with water, the booms retracted, and again pressurized to 15 P.S.1.G.
- 14. The vibration mechanism is started and readings taken. Adjustments are made as necessary to produce the required vibration.
- 15. Rotation speed of the eccentric weights is adjusted to 2000 +0 -60
- 16. Vibration is stopped at the end of 10 minutes and the tank is depressurized and drained.

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TEST RESULTS:

See page 4.7

RECOMMENDATIONS:

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# RESULTS OF TEST

After 17-1/2 hours of simultaneous slosh and vibration a crack in the outer skin located immediately aft of the aft hook was noted. The test was discontinued at this point and the test assembly was removed from the fixture. Close examination revealed the crack to be approximately 18 inches in circumferential length, 15° in both directions from the top centerline of the assembly. The crack appeared to have started in the smallest radius of the bend adjacent to the "flat" for the aft mounting hook. (Ref. photo on page 4.10 of this report)

When the tank was disassembled and the inner fiberglass tank inspected, the following discrepancies were noted:

- 1. Inner rubber liner loose from tank, torn in several places.
- Several layers of apparently non-impregnated fiberglass strands adhering to the loose rubber liner, also hanging loose in the tank.
- Slosh baffle bulkheads integral but loose and lying on bottom of tank.
- 4. Circumferential ring to hold bulkheads still intact.
- Small partial ring segments to retain bulkheads entirely loose.
- 6. The exposed inside surfaces of the inner tank appeared to be smooth.

The photograph on page 4.11 shows the condition of the inner tank immediately after removal from test fixture.

The outer skin of the tank was repaired in accordance with the sketch on page 4.12. The slosh baffles and rubber liner were removed from the inner tank, and the assembly was again subjected to the complete test. The tank then successfully completed the test in accordance with the procedure previously outlined.

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APPROVED			General mills tank assy	REPOR		

## RESULTS OF TEST (contd)

A displacement of .038 inches double amplitude at the tips of the booms were noted during the 15 minutes of slosh-vibration with the booms extended. At the conclusion of the test the booms were again extended and a scan for resonance was made with the rollowing results:

- 1. Resonant frequency of booms 1650 C.P.M.
- Displacement at resonant frequency .35 inch double amplitude at tips.

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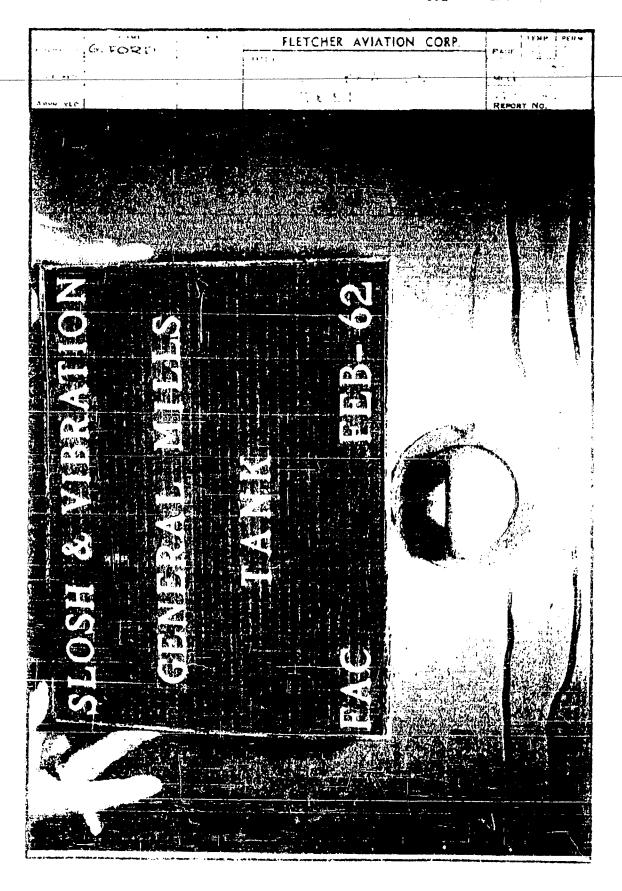
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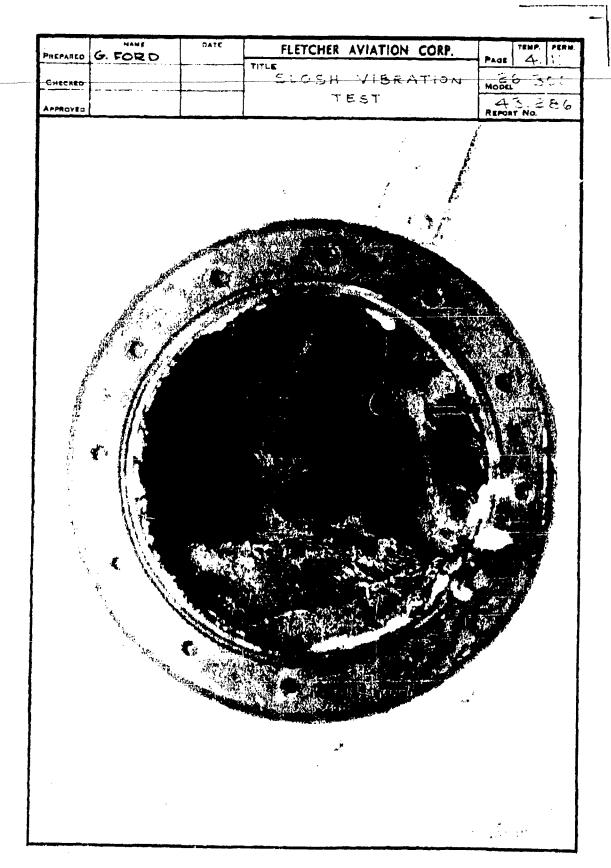
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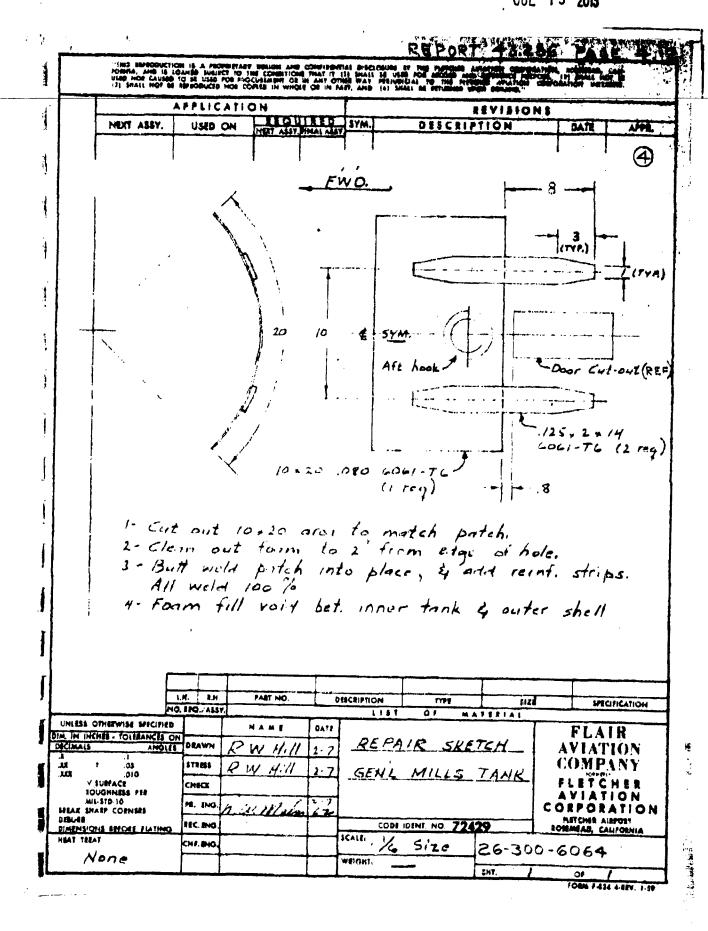
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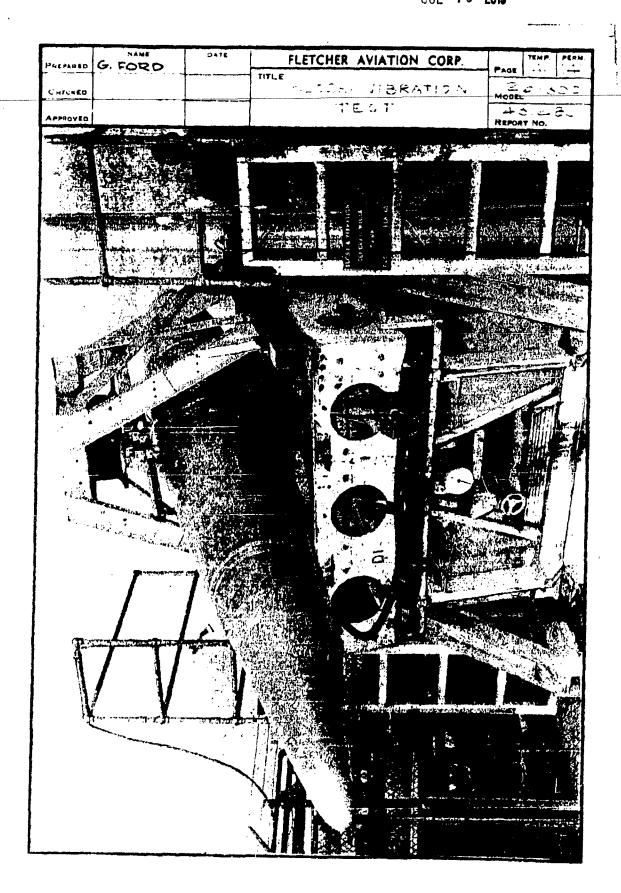


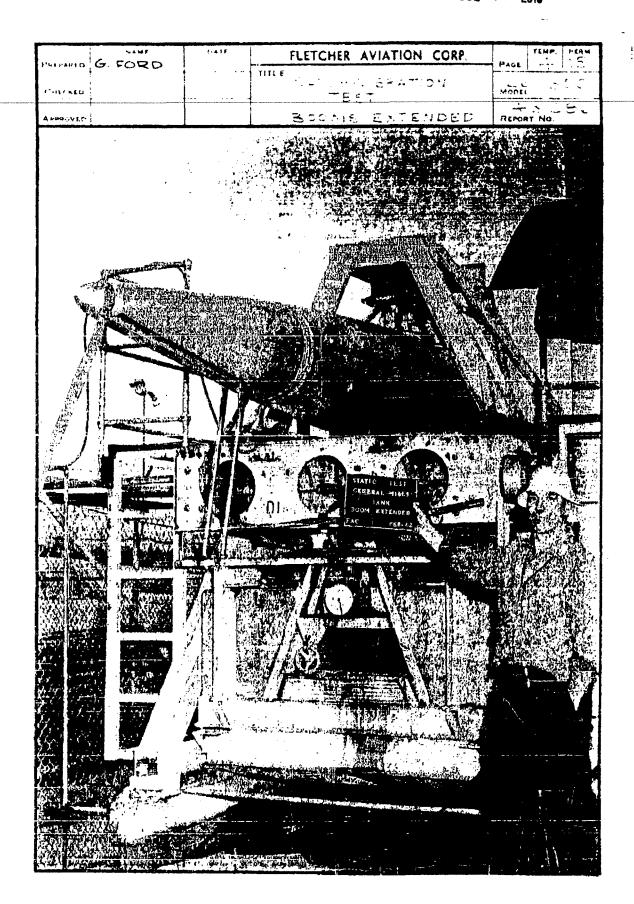
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G. FORD FLETCHER AVIATION CORP. TITLE E E REPORT NO.





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PREPARED W	. Callahan	1-5-62	LEGICULE WHELLOW COMPANY	PAGE		5.0
	A. IL SPANIS MAN	معاكات مرجع ليست كيب	TITLE			
CHECKED			LEAKAGE TEST	MODEL	. 26-	300
		) <del></del>	GENERAL MILLS TANK ASSY		43.	286
APPROVED :		,		REPOR	T No.	

#### ADMINISTRATIVE DATA

(3)

## PURPOSE OF TEST:

To demonstrate that the tank will withstand the required internal pressure without leakage.

## MANUPACTURER:

Pletcher Aviation Company

MANUFACTURER'S MODEL NO.:

26-300

ASSEMBLY DRAWING

26-300-48031

QUANTITY OF ITEMS:

One (1)

SECURITY CLASSIFICATION:

None

TEST DATE:

2-12-62

TEST CONDUCTED BY:

Wayne Callahan

DISPOSITION OF SPECIMEN:

Use for ejection test.

#### ABSTRACT:

The assembly successfully met all of the requirements of the test.

PREPARED	W.Callahan	1-5-62	FLETCHER AVIATION COMPANY	PAGE	tesip.	768B.
CHECKED			LEAKAGE TEST	MODEL	26-	300
APPROVED			General Mills Tank assy	Repos	43.	286

#### PACTUAL DATA

#### REQUIREMENTS:

 The inner fiberglass tank with all openings sealed is subjected to an internal pressure of 20 P.S.I.G. using Freon 12. The tank is then checked for leakage using a General Electric H-l tester.

## TEST BOULPMENT:

- 1. Tank Support
- 2. Pressure dage
- 3. Preon 12 container with regulator.
- 4. General Electric H-1 tester.

#### PROCEDURE:

- With the tank supported, a regulated Preon 12 supply line is connected to the assembly.
- 2. A pressure gage is connected to the assembly.
- 3. From 12 pressure is admitted into the assembly until the pressure gage indicates 20 P.S.I.G.
- 4. External surfaces of the assembly are checked for leakage using the General Electric H-1 tester at a sniffing rate of 1/2 inch per second.
- 5. Inspect the tank visually for evidence of failure such as damage to the bulkheads or liner of the inner tank or structural damage to the inner or outer tank.

FAC fem Fact

PREPARED	W.Callulian	DATE 1-5-62	Fletcher aviation company	PAGE	71MF. P18M.
CHECKED		and Section to the second of the	LEAKAGE TEST GENERAL MILLS TANK ASSY	Model	26-300 43.286
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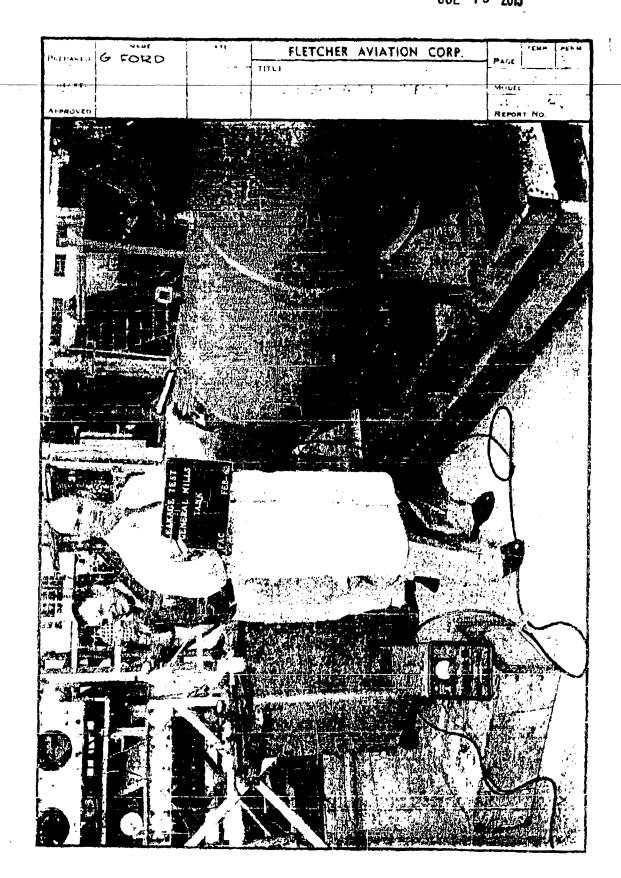
#### TEST RESULTS:

During the leakage test it was determined that the forming process used to install the form between the inner tank and outer shell contained halogen gas. Since the General Electric tester used in probing for leaks is sensitive to this gas, there was a constant leakage indication during the test.

The sensitivity of the instrument was adjusted so that it would not indicate leakage from the presence of gas in the foam. It was then determined that it would indicate leakage if any of the stronger mixture from the inner tank escaped to atmosphere.

The test was concluded with the instrument adjusted per the preceding paragraph, and the test conducted per the outlined procedure on page 5.1. There was no evidence of leakage. No evidence of leakage when pressurized to 20 p.s.i.g.

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CHECKED Approved	,		GROUND EJECTION TEST GROUND EJECTION TEST	Model	43.	300

# 3

## ADMINISTRATIVE DATA

# PURPOSE OF TEST

To determine ejection characteristics.

#### MANUPACTURER

Fletcher Aviation Company

MANUFACTURER'S MODEL NO.

26-300

#### ASSEMBLY DRAWING

26-300-48031

## QUANTITY OF ITEMS

One (1)

## SECURITY CLASSIFICATION

None

## DATE TEST COMPLETED

2-16-62

#### TEST CONDUCTED BY

Wayne Callahan

# DISPOSITION OF SPECIMEN

Hold at Fletcher Aviation Company for sixty (60) days for G.M.I. disposition.

#### **ABSTRACT**

The asacmbly successfully met all of the requirements of the test.

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APPROVED			GENERAL MILLS TANK ASSY	REPOR	43.	286

## PACTUAL DATA

## REQUIREMENTS

That one (1) or more ejection be made with a lightweight tank and one (1) or more ejection be made with heavy-weight tank so as to accurately determine the peak force, velocity, acceleration, and tank attitude at end of stroke. The peak ejection force shall not axceed 30,000 lbs.

## TEST BOULDMENT

- 1. Ejection frame
- 2. Suspension fixture
- 3. Pylon
- 4. Lightweight Store
- 5. Heavyweight Store
- 6. 28V D.C. power supply
- 7. 20,000 Ohm/Volt Multimeter
- 8. Midwestern Oscillograph
- 9. F.A.C. Force Transducer
- 10. Extensometer (Century Eng.)
- 11. Accelerometer
- 12. Amplifier (Miller)
- 13. High Speed Camera (Wollensak)
- 14. Goose Control

## TEST PROCEDURE

- 1. Install suspension fixture on ejection frame.
- 2. Install pylon on suspension fixture.
- 3. Install lightweight store on pylon.
- 4. Apply 28V D.C. to the mechanism.
- 5. Measure voltage at firing pins.
- 6. Remove safety pin.
- Measure voltage at firing pins.
- 8. Disconnect power supply and install safety pin.

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			GROUND EJECTION TEST		26-300
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l			General Mills Takk assy	1 .	43.286
APPROVED	<u> </u>			REPOR	it No.

## PACTURE DATA

## TEST PROCEDURE (contd)

- 11. Install extensometer.
- 12. Install camera.
- 13. Check instrument and camera circuit.
- 14. Install cartridges.
- 15. Remove safety pin.
- 16. Activate instrument and camera circuits.
- 17. Activate firing mechanism.
- 18. Clean pylon assembly and inspect.
- 19. Install heavyweight store on pylon.
- 20. Repeat steps 4 through 17.

FLETCHER AVIATION COMPANY PREPARED W. Callahan 1-8-62 TITLE GROUND EJECTION TEST Moost 26-300 GENERAL MILLS TANK ASSY 43,286 APPROVED REPORT NO. 3 TEST RESULTS Lightweight Heavyweight Tank Tent 15,600 lbs. 21.956 lbs. 37.1 "G's" Peak Ejection Force Peak Acceleration 27.4 "G'8" Peak Velocity 21.9 ft/sec. 17.8 ft/sec. Attitude at end of stroke Level RECOMMENDATIONS: F.A.C. ENG'R: QUALITY CONTROL: CUSTOMER:

FAC form F108

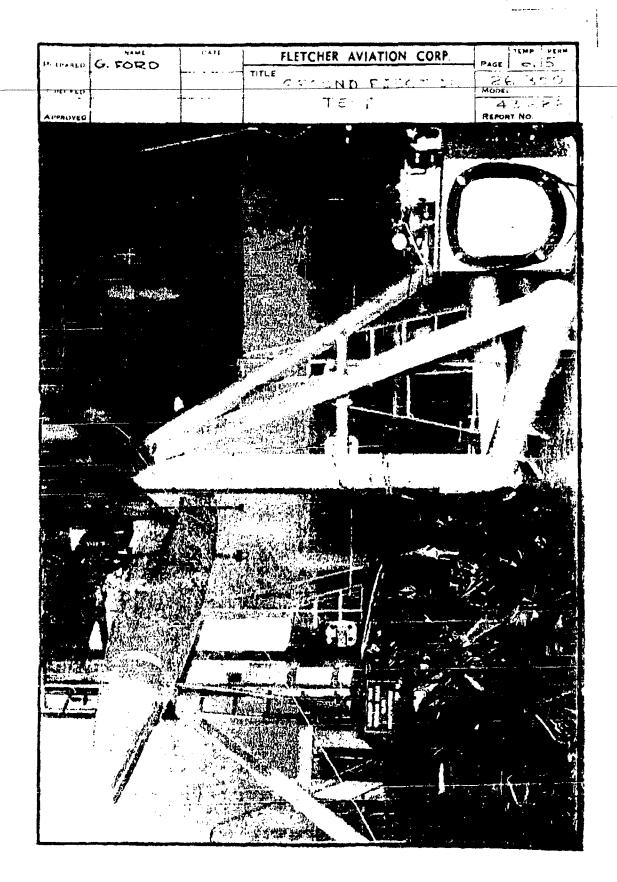
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-DATE 4. FLETCHER AVIATION CORP. G. FORD PREPARED TITLE 26-300 NODEL GROUND EJECTION CHECKED 43.286 TEST APPROVED

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Reviewed Chief, RDD, WHS IAW EO 13526, Section 3.5 Date:

JUL 19 2013



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PREPARED	W.Callahan	1-6-62	FLETCHER AVIATION COMPANY	PAGE	TÉMA.	7.0
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APPROVED		:		REFOR	T NO.	

#### ADMINISTRATIVE DATA

3

#### PURPOSE OF TEST:

To demonstrate the structural integrity of the tank.

## MANUPACTURER:

Fletcher Aviation Company

MANUFACTURER'S MODEL NO.

26-300

#### ASSEMBLY DRAWING:

26-300-

## QUARTITY OF ITEMS:

One (1)

# SECURITY CLASSIFICATION:

None

## TEST DATE:

2-8-62 through 2-12-62

## TEST CONDUCTED BY:

Wayne Callahan

# DISPOSITION OF SPECIMEN:

Hold at F.A.C. for 60 days for g.M.I. disposition.

#### ABSTRACT:

The assembly successfully met all of the requirements of the test.

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## PACTUAL DATA

# **(1)**

#### TEST REQUIREMENT

The tank is subjected to the following test conditions and does support these conditions without failure.

A tank that has not been used for other tests shall be used for this test. The static test tank shall be a complete structure, less such non-structural elements as turbine generator, booms, actuator, boom support structure, fluid handling components, and electrical components. Also, the inner fiberglass tank shall be simulated by a suitable structure of the same size and configuration. The static test tank shall be of the same quality workmanship as the flight tank delivered on the contract and shall be, except for the inner tank, structurally identical to the flight tank as indicated in the reports and drawings submitted.

The test techniques of the tank is as follows:

The tank support jig shall be constructed to duplicate the attach point locations of the Aero 7A rack to produce the most critical hook and sway brace reactions.

Loads are introduced into the test tank by means of tension pads internal formers, and/or external straps. Of these methods, tension pads are preferable in that they allow a better load distribution and are less susceptable to local ovarloading difficulties. Care is taken to insure that the load application devices do not materially affect the strength of the test tank by introducing artificial stiffness, etc.

All applied test loads are suitably monitored by calibrated equipment (pressure gages, load dynamometers, etc.) so that acceptable test accuracy is obtained.

Internal pressure, where applicable, is applied pneumatically or hydraulically and the pressure suitably monitored with calibrated measuring devices.

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## FACTUAL DATA (contd)

Tare weight of store and all load application devices is accounted for in all test loadings. Particular care is taken when dense fluids (water, stc.) are used for introducing internal pressure so that true incremental loads for all load components are obtained. Independent application of combined load components (that is vertical, side, and aft loads) are preferable over resultant load application to facilitate maintaining correct relation—ship of load components with each other for full range of load from zero to ultimate.

JUL 19 2013

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The test loads of the tank are as specified in contract.

# TEST POULPMENT:

- 1. Static test frame.
- 2. Tank support fixture.
- 3. Loading pads, whiffle trees, etc., per applicable test conditions.
- 4. Hydraulic pumps.
- 5. Hydraulic cylinders (jacks) with net areas as listed on test data sheets.
- 6. Hydraulic test gages: as necessary
- 7. Pour 24-inch Sturrett steel engine marked scales reading to .010 inch.
- 8. Surveyor's level.

#### TEST PROCEDURE:

All test conditions are run with the test tank mounted in the horizontal position in the test jig. The test procedure is identical in each case and consists of the following steps:

- The system of loading jacks and whiffle trees is installed, checked functionally, and inspected for proper location.
- Readings of deflection at zero load is taken by means of a series of steel scales hung along the length of the tank, and a surveyor's level. Lateral deflections are measured from a wire stretched alongside of the tank.
- 3. The load is then applied in increments of 25% of limit load, and the deflection readings taken at 25, 50, 75, and 100%
- 4. The jack loads are then reduced to zero, and deflection readings taken to check possible permanent set.
- 5. Load is again applied in 25% increments up to 100% of limit load.

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# TEST PROCEDURE: (CONT.)

- 6. The static load is increased to 125% of limit load.
- 7. Deflection readings are taken.
- 8. The static load is increased to 150% of limit load, and deflection readings are taken. (150% of limit load = 100% ultimate load.)
- 9. The jack loads are reduced to zero and deflection readings are taken.

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APPROVED		}	GENERAL MILLS TANK	Repor	43.	286

#### STRUCTURAL STATIC TESTS

A review of F.A.C. Report No. 43.284, "Loads and Stress Analysis, General Mills Tank", shows that there are three design conditions critical for the tank and its attaching structure. These are conditions #10, 25, and 29. Condition #29 is an ejection condition which will be adequately covered by the actual tank full and empty ejections. (Ref. pages 6.0 to 6.3 incl.) Test loads for Condition #10 and 25 are developed on the following pages.

As on page #2 of Report #43.284, the sign convention for the test loads is as follows:

2 = Upward acting

Y = Acting to the left

X " Aft acting

Positive moment vectors are in the same direction using the left-hand rule.

The unit loads shown on page 7.5 are revised to agree with the latest weight data in Appendix "A" of Report #43.284.

All load factors, loads and moments in this section, are on an ultimate basis.  $(1-1/2 \times 1)$  init load. The tank itself will be tested empty and unpressurized.

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APPROVED			Static Tests	Repor	43.2 11 No.	84

# UNIT INERTIA LOAD DISTRIBUTION (Revised per Appendix

	Tank	Full	1 Full,	Aft CG.
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Test Logits - Cond. "25 (Tank Full)

 $I_{yy} = I_{\frac{2}{2}} = 772.6 \text{ s/ug-ft}^{2} (Pg. ^{4}2, rep. ^{4}3.284)$   $M_{yy} = 12 \times 772.6 \times (-18.00) = -166,880 \text{ in-16s}$   $M_{77} = 12 \times 772.6 \times 7.00 = 83.440 \text{ in-16s}$ 

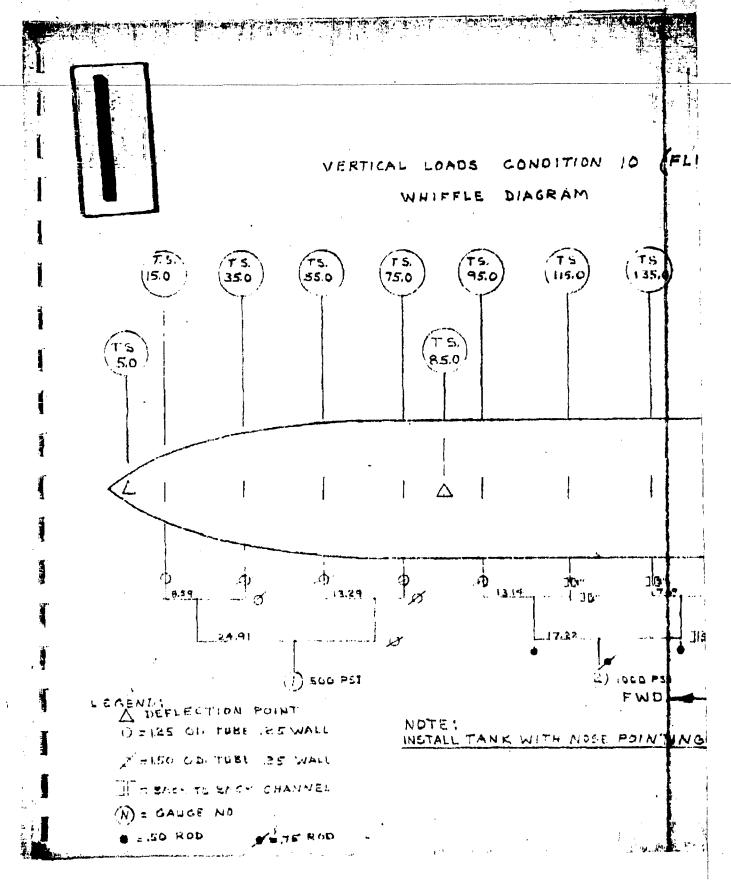
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Reviewed Chief, RDD, WHS IAW EO 13526, Section 3.5 Date:

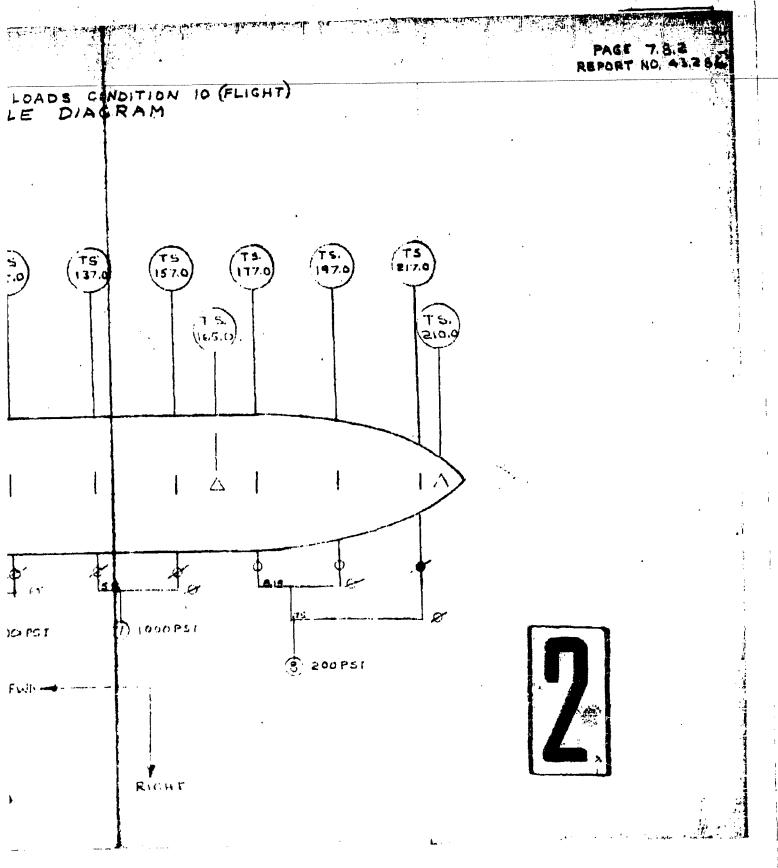
JUL 19 2013

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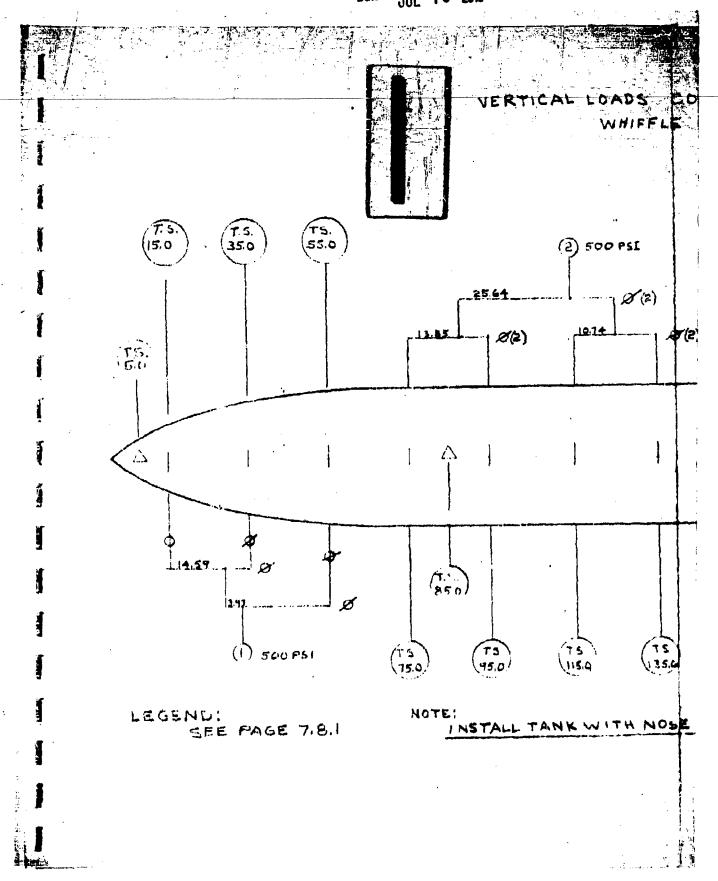
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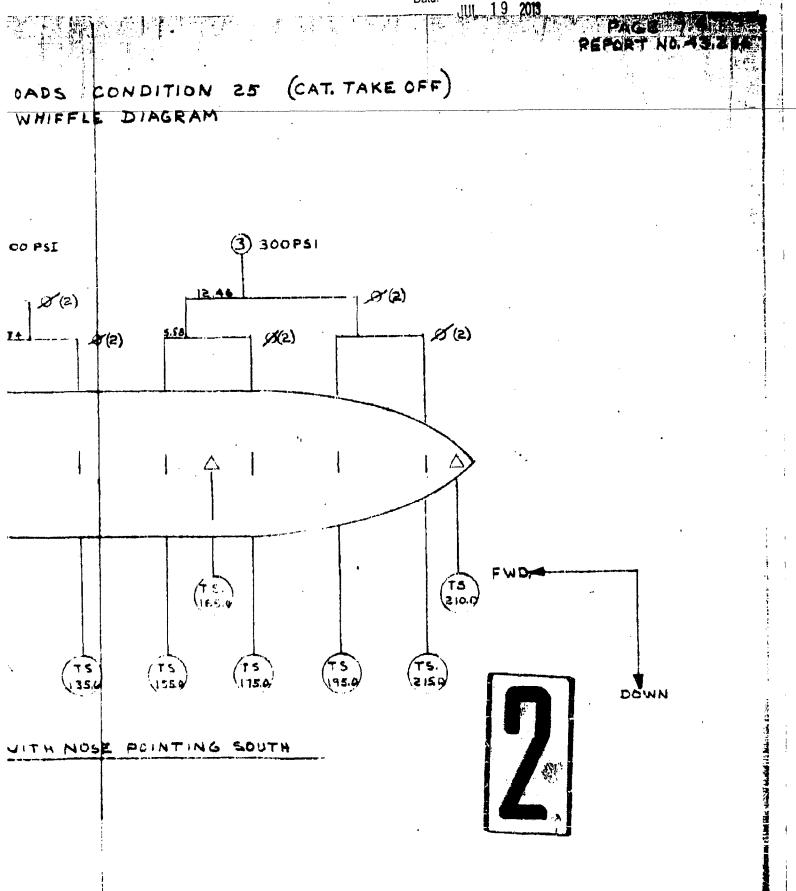
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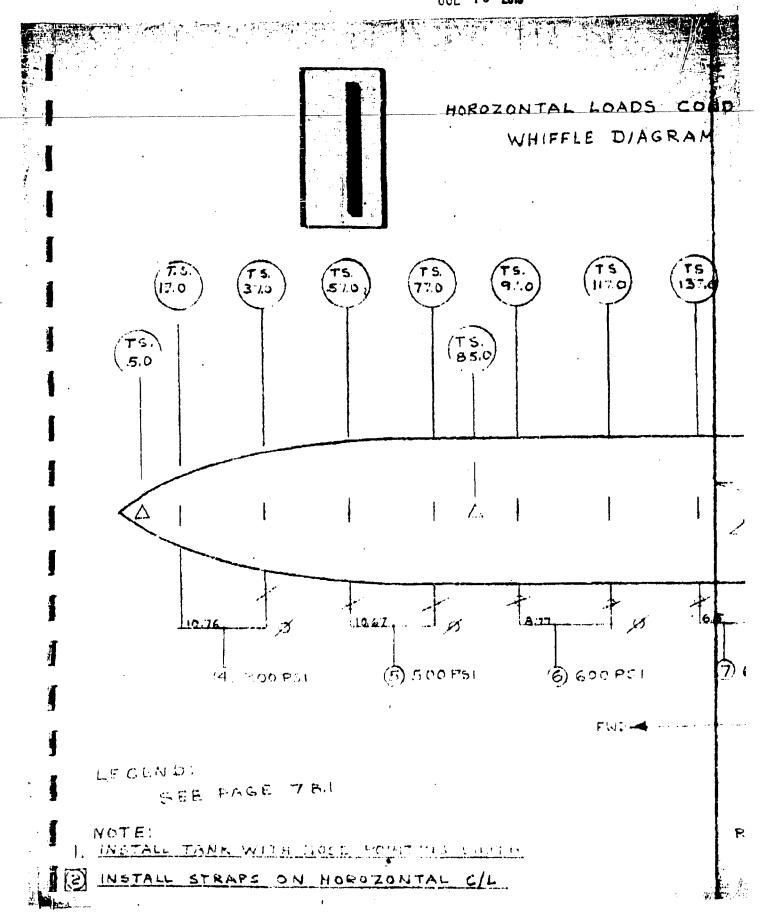


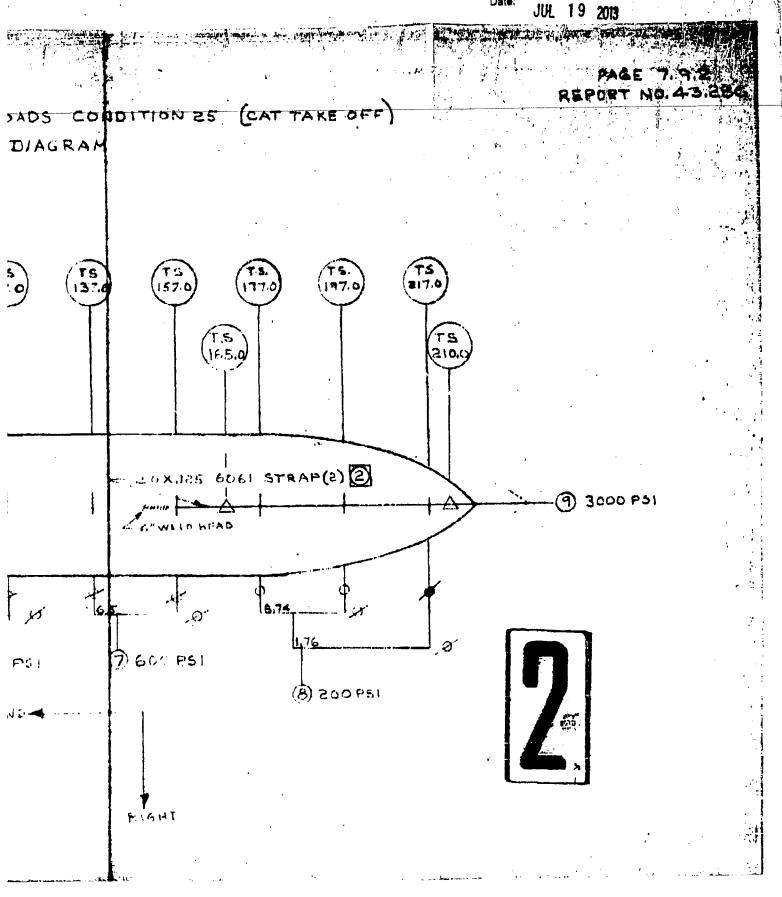
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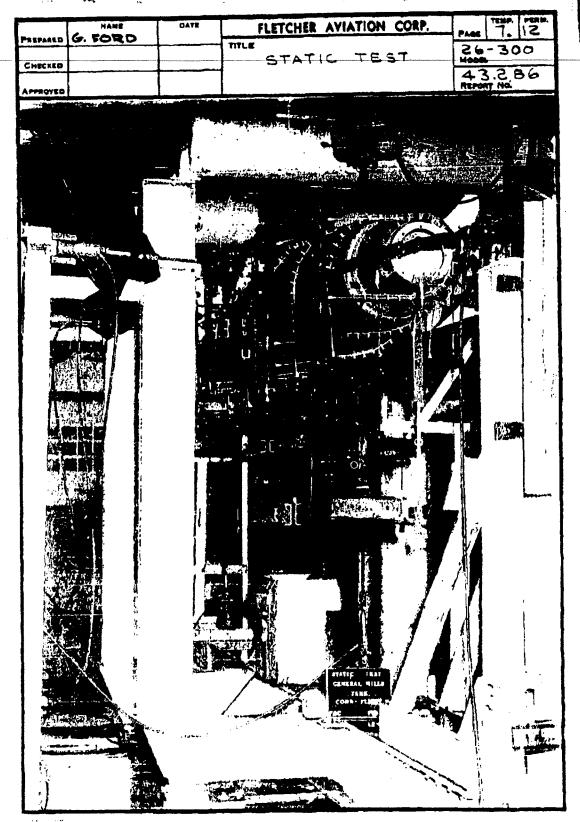




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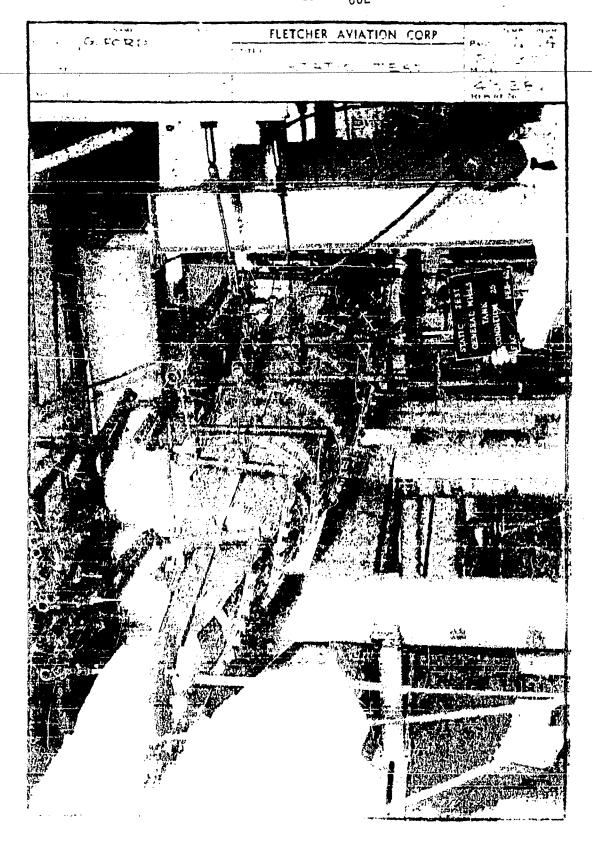
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JUL 19 2013



- Address







## DEPARTMENT OF DEFENSE WASHINGTON HEADQUARTERS SERVICES

1155 DEFENSE PENTAGON WASHINGTON, DC 20301-1155



MEMORANDUM FOR DEFENSE TECHNICAL INFORMATION CENTER
(ATTN: WILLIAM B. BUSH)
8725 JOHN J. KINGMAN ROAD, STE 0944

AUG 1 2013

FT. BELVIOR, VA 22060-6218

SUBJECT: OSD MDR Cases 12-M-3144 through 12-M-3156

At the request of the documents, we have conducted a Mandatory Declassification Review of the documents in the above referenced cases on the attached Compact Disc (CD) under the provisions of Executive Order 13526, section 3.5, for public release. We have declassified the documents in full. We have attached a copy of our response to the requester. If you have any questions, please contact Ms. Luz Ortiz by phone at 571-372-0478 or by e-mail at luz.ortiz@whs.mil, luz.ortiz@osd.smil.mil, or luz.ortiz@osdj.ic.gov.

Robert Storer Chief, Records and Declassification Division

Policet Storen

## Attachments:

- 1. MDR request w/ document list
- 2. OSD response letter
- 3. CD (U)



April 26, 2012

Department of Defense
Directorate for Freedom of Information and Security Review
Room 2C757
1155 Defense Pentagon
Washington, D.C. 20301-1155

Sir:

I am requesting under the Mandatory Declassification Review provisions of Executive Order 13291, copies of the following documents. I have tried several times to acquire them through DTIC, but the sites stated they are not available.

I am conducting research into the previous methods used to disseminate biological agents. Many source I use to have access to have been deleted from the internet. On numerous occasions I have been informed that formerly classified information that was declassified, have now become classified again (since 911). My attempts to locate such Executive Orders, regulations, laws, or other changes to this question have not successful nor revealed a specific source. As such I would appreciate any information you can shed on this question.

## Documents requested.

AD 348405, Dissemination of Solid and Liquid BW (Biological Warfare) Agents Quarterly 12-M-3144 Progress Report Number 14, 4 Sept - 4 Dec 1963, G. R. Whitnah, February 1964, General Mills Report number 2512, General Mills, Inc., Minneapolis, MN, Contract number DA 18064 CML 2745,102 pages. Prepared for U.S. Army Biological Laboratories, Fort Detrick, Maryland. Approved by S.P. Jones, Director of Aerospace Research at General Mills. Project No. 82408. General Mills Aerospace Research Division, 2295 Walnut Street, St. Paul 13, Minnesota. AD 346751, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3145 Progress Report Number 12, March 4 - June 4, 1963, G. R. Whitnah, July 1963, General Mills Report number 2411, General Mills, Inc., Minneapolis, MN, Contract number DA 18064 CML 2745. 184 pages. Approved by S.P. Jones, Director of Aerospace Research at General Mills. Project No. 82408. General Mills Aerospace Research Division, 2295 Walnut Street, St. Paul 13, Minnesota.

AD 346750, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3146 Progress Report Number 13, 4 June - 4 Sept 1962, G.R. Whitnah, October 1963, General Mills

Report number 2451, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 19 pages (?)

AD 332404, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3147 Progress Report Number 7, Dec. 4, 1961 - March 4, 1962, by G.R. Whitnah, February 1963, General Mills Report Number 2373, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 123 pages.

AD 333298, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-314 B Progress Report Number 9, June 4, 1962 - Sept. 4, 1962. by G.R. Whitnah, October 1962, General Mills Report Number 2344, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 130 (or 150) pages.

AD 332405, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3147 Progress Report Number 8, Period March 4, 1962 - June 4, 1962. G.R. Whitnah, August 1962, General Mills Report Number 2322, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 198 pages.

AD 329067, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-W-3150 Progress Report Number Six, G.R. Whitnah, February 1962, General Mills Report Number 2264, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 103 pages. Approved by S.P. Jones, Manager, Materials and Mechanics Research, General Mills Research and Development Office, 2003 East Hennepin Avenue, Minneapolis 13, Minnesota.

AD 327072, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly /2-M-3157 Progress Report Number Five, 4 June - 4 Sept 1961. by G.R. Whitnah, November 1961, General Mills Report Number 2249, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745.

AD 325247, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3152 Progress Report Number 4, 4 March - 4 June 1961, by J.E. Upton for G.R. Whitnah, Project Manager. February 1963, General Mills Report Number 2216, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. General Mills Electronics Group, Research Dept., 2003 East Hennepin Avenue, Minneapolis 13, Minnesota. 225 pages.

AD 324746, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Progress 12-M-3133 Report 3 Juen - 3 Sept. 1960. by G.R. Whitnah, October 1960, General Mills Report Number 2125, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 78 pages

AD 323599, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3154 Progress Report Number 2, for period 4 Sept - 4 Dec 1960, by G.R. Whitnah, February 1961, General Mills Report Number 2161, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 90 pages? Mechanical Division of General Mills, Inc., Research Department, 2003 East Hennepin Avenue, Minneapolis 13, Minnesota.

AD 323598, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3175 Progress Report, for period 4 Dec. 1960 - 4 March 1961, by G.R. Whitnah, May 1961, General Mills Report Number 2200, General Mills, Inc., Minneapolis, MN, Contract Number DA 18064 CML 2745. 95 pages.

AD 337635, Dissemination of Solid and Liquid BW (Biological Warfare) Agents, Quarterly 12-M-3156 Progress Report No. 10, period Sept. 4, 1962 - Dec. 4, 1962. G.R. Whitnah, Project Manager, Approved by S.P. Jones, Aerospace Research, February 1963. 247 pages.

## Sincerely

